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(54) Title: DEVELOPMENT OF NOVEL ANTI-MICROBIAL AGENTS BASED ON BACTERIOPHAGE GENOMICS (57) Abstract A method for identifying suitable targets for antibacterial agents based on identifying targets of bacteriophage-encoded proteins is described. Also described are compositions useful in the identification methods and in inhibiting bacterial growth, and methods for preparing and using such compositions.			

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DESCRIPTION

Development of Novel Anti-Microbial Agents Based on Bacteriophage Genomics

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BACKGROUND OF THE INVENTION

The present invention relates to the field of antibacterial agents and the treatment of infections of animals or other complex organisms by bacteria.

10 The frequency and spectrum of antibiotic-resistant infections have, in recent years, increased in both the hospital and community. Certain infections have become essentially untreatable and are growing to epidemic proportions in the developing world as well as in institutional settings in the developed world. The staggering spread of antibiotic resistance in pathogenic bacteria has been attributed to microbial
15 genetic characteristics, widespread use of antibiotic drugs, and changes in society that enhance the transmission of drug-resistant organisms. This spread of drug resistant microbes is leading to ever increasing morbidity, mortality and health-care costs.

Ironically, it is the very success of antibiotics, resulting in their widespread use, that has contributed the most to rising numbers of drug resistant bacterial strains.
20 The longer a bacterial strain is exposed to a drug, the more likely it is to acquire resistance. Today, a total of 160 antibiotics, all based on a few basic chemical structures and targeting a small number of metabolic pathways, have found their way to market. Over-prescription of these drugs, as well as the failure of patients to comply with the complete antibiotic regimen, has lead to the rapid emergence of
25 antibiotic resistant strains. Such misuse of prescriptions, careless use of antibiotics in virtually all commercial production of beef and fowl, and changing societal conditions, such as the growth of day-care centers, increased long-term care in hospitals, and increased mobility of the population, has provided an environment where drug-resistant microbes can emerge and spread. Thus, virtually all common
30 infectious bacteria are becoming, or have already become, resistant to one or more groups of antibiotics. Such resistance now reaches all classes of antibiotics currently in use, including: β -lactams, fluoroquinolones, aminoglycosides, macrolide peptides, chloramphenicol, tetracyclines, rifampicin, folate inhibitors, glycopeptides, and mupirocin.

35 Over the last 45 years bacteria have adapted genetically to avoid the destruction/alteration of the essential pathways that these chemotherapeutic agents

target. Antibiotic resistant bacterial strains are now emerging at a higher rate than the rate at which new antibiotics are being developed. The consequence of this dilemma has been a dramatic increase in the cost of treating infections what would otherwise easily succumb to routine antibiotic therapy. Furthermore, and perhaps most
5 importantly, the emergence of multiple drug resistant pathogenic bacteria has led to a significant increase in morbidity and mortality, particularly in institutional settings.

Most major pharmaceutical companies have on-going drug discovery programs for novel anti-microbials. These are based on screens for small molecule inhibitors (natural products, bacterial culture media, libraries of small molecules,
10 combinatorial chemistry) of crucial metabolic pathways of the micro-organism of interest (*e.g.*, bacteria, fungi, parasites, worms). The screening process is largely for cytotoxic compounds and in most cases is not based on a known mechanism of action of the compounds. Pharmaceutical companies have large programs in this area. Classical drug screening programs are being exhausted and many of these
15 pharmaceutical companies are looking towards rational drug design programs.

Several small to mid-size biotechnology companies as well as large pharmaceutical companies have developed systematic high-throughput sequencing programs to decipher the genetic code of specific micro-organisms of interest. The goal is to identify, through sequencing, unique biochemical pathways or intermediates
20 that are unique to the microorganism. Knowledge of this may, in turn, form the rationale for a drug discovery program based on the mechanism of action of the identified enzymes/proteins. Genome Therapeutics Corp., The Institute for Genome Research, Human Genome Sciences Inc., and other companies have such sequencing programs in place. However, one of the most critical steps in this approach is the
25 ascertainment that the identified proteins and biochemical pathways are 1) non-redundant and essential for bacterial survival, and 2) constitute suitable and accessible targets for drug discovery.

SUMMARY OF THE INVENTION

While animals such as humans are, on occasion, infected by pathogenic bacteria, bacteria also have natural enemies. A number of host-specific viruses, known as bacteriophages or phages, infect and kill bacteria in the natural environment. Such bacteriophages generally have small compact genomes and bacteria are their exclusive hosts. Many known bacteria are host to a large number of bacteriophages that have been described in the literature. During the 1940's - 1960's, phage biology was an area of active research. As a testimony to this, the study of phages which infect and inhibit the enteric bacterium *Escherichia coli* (*E. coli*) contributed much to the early understanding of molecular biology and virology.

As is generally understood, bacteriophage (or phages) are viruses that infect and kill bacteria. They are natural enemies of bacteria and, over the course of evolution, have developed proteins (products of DNA sequences) which enable them to infect a host bacteria, replicate their genetic material, usurp host metabolism, and ultimately kill their host. The scientific literature well documents the fact that many known bacteria have a large number of such bacteriophages (Ackermann and DuBow, 1987) that can infect and kill them (for example, see the ATCC bacteriophage collection at <http://www.atcc.org>).

This invention utilizes the observation that bacteriophages successfully infect and inhibit or kill host bacteria, targeting a variety of normal host metabolic and physiological traits, some of which are shared by all bacteria, pathogenic and nonpathogenic alike. The term "pathogenic" as used herein denotes a contribution to or implication in disease or a morbid state of an infected organism. The invention thus involves identifying and elucidating the molecular mechanisms by which phages interfere with host bacterial metabolism, an objective being to provide novel targets for drug design. Whether the phage blocks bacterial RNA transcription or translation, or attacks other important metabolic pathways, such as cell wall assembly or membrane integrity, the basic blueprint for a phage's bacteria-inhibiting ability is encoded in its genome and can be unlocked using bioinformatics, functional genomics, and proteomics. By these means, the invention utilizes sequence information from the genomics of bacteriophage to identify novel antimicrobials that can be further used to actively and/or prophylactically treat bacterial infection.

Two important components of the invention thus are: i) the identification of bacteria-inhibiting phage open reading frames ("ORF"s) and corresponding products that can be used to develop antibiotics based on amino acid sequence and secondary structural characteristics of the ORF products, and ii) the use of bacteriophages to map

out essential bacterial target genes and homologs, which can in turn lead to the development of suitable anti-microbial agents. These two avenues represent new and general methods for developing novel antimicrobials.

The invention thus concerns the identification of bacteriophage ORFs that supply bacteria-inhibiting functions. In this regard, use of the terms "inhibit", "inhibition", "inhibitory", and "inhibitor" all refer to a function of reducing a biological activity or function. Such reduction in activity or function can, for example, be in connection with a cellular component, *e.g.*, an enzyme, or in connection with a cellular process, *e.g.*, synthesis of a particular protein, or in connection with an overall process of a cell, *e.g.*, cell growth. In reference to bacterial cell growth, for example, an inhibitory effect (*i.e.*, a bacteria-inhibiting effect) may be bacteriocidal (killing of bacterial cells) or bacteriostatic (*i.e.*, stopping or at least slowing bacterial cell growth). The latter slows or prevents cell growth such that fewer cells of the strain are produced relative to uninhibited cells over a given period of time. From a molecular standpoint, such inhibition may equate with a reduction in the level of, or elimination of, the transcription and/or translation of a specific bacterial target(s), or reduction or elimination of activity of a particular target biomolecule.

It is particularly advantageous to evaluate a plurality of different phage ORFs for inhibitory activity that may be from one, but is preferably from a plurality of different phage. For example, evaluating ORFs from a number of different phage of the same bacterial host provides at least two advantages. One is that the multiple phages will provide identification of a variety of different targets. Second, it is likely that multiple phage will utilize the same cellular target

As used herein, the terms "bacteriophage" and "phage" are used interchangeably to refer to a virus which can infect a bacterial strain or a number of different bacterial strains.

In the context of this invention, the term "bacteriophage ORF" or "phage ORF" or similar term refers to a nucleotide sequence in or from a bacteriophage. In connection with a particular ORF, the terms refer an open reading frame which has at least 95% sequence identity, preferably at least 97% sequence identity, more preferably at least 98% sequence identity with an ORF from the particular phage identified herein (*e.g.*, with an ORF as identified herein) or to a nucleic acid sequence which has the specified sequence identify percentage with such an ORF sequence.

A first aspect of the invention thus provides a method for identifying a bacteriophage nucleic acid coding region encoding a product active on an essential bacterial target by identifying a nucleic acid sequence encoding a gene product which

provides a bacteria-inhibiting function when the bacteriophage infects a host bacterium, preferably one that is an animal or plant pathogen, more preferably a bird or mammalian pathogen, and most preferably a human pathogen. The bacteriophage is an uncharacterized bacteriophage. Thus, the method excludes, for example, phage
5 λ , ϕ x174, m13 and other *E.coli*-specific bacteriophage that have been studied with respect to gene number and/or function. It also excludes, for example, the nucleic acid coding regions described in Tables 12-14, and in preferred embodiments, excludes the phage in which those regions are naturally located.

In connection with bacteriophage, the term "uncharacterized" means that a
10 certain bacteriophage's genome has not yet been fully identified such that the genes having function involved in inhibiting host cells have not been identified. In particular, phage for which the description of genomic or protein sequence was first provided herein are uncharacterized. Phage sequences for which host bacteria-inhibiting functions have been identified prior to the filing of the present application
15 (or alternatively prior to the present invention) are specifically excluded from the aspects involving utilization of sequences from uncharacterized bacteriophage, except that aspects may involve a plurality of phage where one or more of those phage are uncharacterized and one or more others have been characterized to some extent. A number of different bacteria-inhibiting phage ORFs are indicated in Tables 11-14.
20 The phage ORFs or sequences identified therein are not within the term "uncharacterized; alternatively, in preferred embodiments the phage containing those ORFs are excluded from this term. Further, any additional phage ORFs (or alternatively the phage which contain those ORFs) which have previously been described in the art as bacteria-inhibiting ORFs are expressly excluded; those ORFs or
25 phage are known to those skilled in the art and the exclusion can be made express by specifically naming such ORFs or phage as needed (likewise for uncharacterized targets as described below). For the sake of brevity, such a listing is not expressly presented, as such information is readily available to those skilled in the art.

Stating that an agent or compound is "active on" a particular cellular target,
30 such as the product of a particular gene, means that the target is an important part of a cellular pathway which includes that target and that the agent acts on that pathway.

Thus, in some cases the agent may act on a component upstream or downstream of the stated target, including on a regulator of that pathway or a component of that pathway.

By "essential", in connection with a gene or gene product, is meant that the host
35 cannot survive without, or is significantly growth compromised, in the absence depletion, or alteration of functional product. An "essential gene" is thus one that encodes a product that is beneficial, or preferably necessary, for cellular growth in

vitro in a medium appropriate for growth of a strain having a wild-type allele corresponding to the particular gene in question. Therefore, if an essential gene is inactivated or inhibited, that cell will grow significantly more slowly, preferably less than 20%, more preferably less than 10%, most preferably less than 5% of the growth rate of the uninhibited wild-type, or not at all, in the growth medium. Preferably, in the absence of activity provided by a product of the gene, the cell will not grow at all or will be non-viable, at least under culture conditions similar to the *in vivo* conditions normally encountered by the bacterial cell during an infection. For example, absence of the biological activity of certain enzymes involved in bacterial cell wall synthesis can result in the lysis of cells under normal osmotic conditions, even though protoplasts can be maintained under controlled osmotic conditions. In the context of the invention, essential genes are generally the preferred targets of antimicrobial agents. Essential genes can encode target molecules directly or can encode a product involved in the production, modification, or maintenance of a target molecule.

A "target" refers to a biomolecule that can be acted on by an exogenous agent, thereby modulating, preferably inhibiting, growth or viability of a cell. In most cases such a target will be a nucleic acid sequence or molecule, or a polypeptide or protein. However, other types of biomolecules can also be targets, *e.g.*, membrane lipids and cell wall structural components.

The term "bacterium" refers to a single bacterial strain, and includes a single cell, and a plurality or population of cells of that strain unless clearly indicated to the contrary. In reference to bacteria or bacteriophage, the term "strain" refers to bacteria or phage having a particular genetic content. The genetic content includes genomic content as well as recombinant vectors. Thus, for example, two otherwise identical bacterial cells would represent different strains if each contained a vector, *e.g.*, a plasmid, with different phage ORF inserts.

In preferred embodiments, the phage is *Staphylococcus aureus* phage 77, 3A, 96, or 44 AHJD, *Enterococcus* sp. phage 182, or *Streptococcus pneumoniae* phage Dp-1.

In preferred embodiments, the phage is selected from. Preferred embodiments involve expressing at least one recombinant phage ORF(s) in a bacterial host followed by inhibition analysis of that host. Inhibition following expression of the phage ORF is indicative that the product of the ORF is active on an essential bacterial target. Such evaluation can be carried out in a variety of different formats, such as on a support matrix such as a solidified medium in a petri dish, or in liquid culture.

Preferably a plurality of phage ORFs are expressed in at least one bacterium. The plurality of phage ORFs can be from one or a plurality of phage. With respect to a single phage or at least one phage in a plurality of phages, the plurality of expressed ORFs preferably represents at least 10%, more preferably at least 20%, 40%, or 60%, still more preferably at least 80% or 90%, and most preferably at least 95% of the ORFs in the phage genome. Preferably, for a plurality of phage, the plurality of expressed ORFs preferably represents at least 10%, more preferably at least 20%, 40%, or 60%, still more preferably at least 80% or 90%, and most preferably at least 95% of the ORFs in the phage genome of each phage. The plurality of phage ORFs can be expressed in a single bacterium, or in a plurality of bacteria where one ORF is expressed in each bacterium, or in a plurality of bacteria where a plurality of ORFs are expressed in at least one or in all of the plurality of bacteria, or combinations of these.

In embodiments of the above aspect (as well as in other aspects herein) in which a plurality of phage are utilized, a plurality of phage have the same bacterial host species; have different bacterial host species; or both. The plurality of phage includes at least two different phage, preferably at least 3, 4, 5, 6, 8, 10, 15, 20, or more different phage. Indeed, more preferably, the plurality of phage will include 50, 75, 100, or more phage. As described herein, the larger number of phage is useful to provide additional target and target evaluation information useful in developing antibacterial agents, for example, by providing identification of a larger range of bacterial targets, and/or providing further indication of the suitability of a particular target (for example, utilization of a target by a number of different unrelated phage can suggest that the target is particularly stable and accessible and effective) and/or can indicate alternate sites on a target which interact with different inhibitors.

Further embodiments involve confirmation of the inhibitor function of the phage ORF, such as by utilizing or incorporating a control(s) designed to confirm the inhibitory nature of the ORF(s) being evaluated. The control can, for example, be provided by expression of an inactive or partially inactive form of the ORF or ORF product, and/or by the absence of expression of the ORF or ORF product in the same or a closely comparable bacterial strain as that used for expression of the test ORF. The reduced level of activity or the absence of active ORF product in the control will thus not provide the inhibition provided by a corresponding inhibitory ORF, or will provide a distinguishably lower level of inhibition. An inactivated or partially inactivated control has a mutation(s), *e.g.*, in the coding region or in flanking regulatory elements, that reduce(s) or eliminate(s) the normal function of the ORF. Thus, the inhibition of a bacterium following expression of a phage ORF is determined by comparison with the effects of expression of an inactivated ORF or the

response of the bacteria in the absence of expression in the same or similar type bacterium. Such determination of inhibition of the bacterium following expression of the ORF is indicative of a bacteria-inhibiting function. These manipulations are routinely understood and accomplished by those of skill in the art using standard techniques. In embodiments utilizing absence of expression of the ORF, the bacteria can, for example, contain an empty vector or a vector which allows expression of an unrelated sequence which is preferably non-inhibitory. Alternatively, the bacteria may have no vector at all. Combinations of such controls or other controls may also be utilized as recognized by those skilled in the art.

10 In embodiments involving expression of a phage ORF in a bacterial strain, in preferred embodiments that expression is inducible.

By "inducible" is meant that expression is absent or occurs at a low level until the occurrence of an appropriate environmental stimulus provides otherwise. For the present invention such induction is preferably controlled by an artificial environmental change, such as by contacting a bacterial strain population with an inducing compound (*i.e.*, an inducer). However, induction could also occur, for example, in response to build-up of a compound produced by the bacteria in the bacterial culture, *e.g.*, in the medium. As uncontrolled or constitutive expression of inhibitory ORFs can severely compromise bacteria to the point of eradication, such expression is therefore undesirable in many cases because it would prevent effective evaluation of the strain and inhibitor being studied. For example, such uncontrolled expression could prevent any growth of the strain following insertion of a recombinant ORF, thus preventing determination of effective transfection or transformation. A controlled or inducible expression is therefore advantageous and is generally provided through the provision of suitable regulatory elements, *e.g.*, promoter/operator sequences that can be conveniently transcriptionally linked to a coding sequence to be evaluated. In most cases, the vector will also contain sequences suitable for efficient replication of the vector in the same or different host cells and/or sequences allowing selection of cells containing the vector, *i.e.*, "selectable markers." Further, preferred vectors include convenient primer sequences flanking the cloning region from which PCR and/or sequencing may be performed.

As knowledge of the nucleotide sequence of phage ORFs is useful, *e.g.*, for assisting in the identification of phage proteins active against essential bacterial host targets, preferred embodiments involve the sequencing of at least a portion of the phage genome in combination with the above methods. This can be done either before or after or independent of expression and inhibition of the ORF in the bacteria, and provides information on the nature and characteristics of the ORF. Such a portion is

preferably at least 10%, 20%, 40%, 80%, 90%; or 100% of the phage genome. For embodiments in which a plurality of phage are utilized, preferably each phage is sequenced to an extent as just specified.

- Such sequencing is preferably accompanied by computer sequence analysis to
- 5 define and evaluate ORF(s), ORF products, structural motifs or functional properties of ORF products, and/or their genetic control elements. Thus, certain embodiments incorporate computer sequence analyses or nucleic acid and/or amino acid sequences. Further, existing data banks can provide phage sequence and product information which can be utilized for analysis and identification of ORFs in the sequence.
- 10 Computer analysis may further employ known homologous sequences from other species that suggest or indicate conserved underlying biochemical function(s) for the inhibitory or potentially inhibitory ORF sequence(s) being evaluated. This can include the sequences of signature motifs of identified classes of inhibitors.

- In the context of the phage nucleic acid sequences, e.g., gene sequences, of this
- 15 invention, the terms "homolog" and "homologous" denote nucleotide sequences from different bacteria or phage strains or species or from other types of organisms that have significantly related nucleotide sequences, and consequently significantly related encoded gene products, preferably having related function. Homologous gene sequences or coding sequences have at least 70% sequence identity (as defined by the
- 20 maximal base match in a computer-generated alignment of two or more nucleic acid sequences) over at least one sequence window of 48 nucleotides, more preferably at least 80 or 85%, still more preferably at least 90%, and most preferably at least 95%. The polypeptide products of homologous genes have at least 35% amino acid
- 25 sequence identity over at least one sequence window of 18 amino acid residues, more preferably at least 40%, still more preferably at least 50% or 60%, and most preferably at least 70%, 80%, or 90%. Preferably, the homologous gene product is also a functional homolog, meaning that the homolog will functionally complement one or more biological activities of the product being compared. For nucleotide or
- 30 amino acid sequence comparisons where a homology is defined by a % sequence identity, the percentage is determined using BLAST programs (with default parameters (Altschul et al., 1997, "Gapped BLAST and PSI-BLAST: a new generation of protein database search programs, Nucleic Acid Res. 25:3389-3402). Any of a variety of algorithms known in the art which provide comparable results can also be used, preferably using default parameters. Performance characteristics for
- 35 three different algorithms in homology searching is described in Salamov et al., 1999, "Combining sensitive database searches with multiple intermediates to detect distant

homologues." *Protein Eng.* 12:95-100. Another exemplary program package is the GCG™ package from the University of Wisconsin.

Homologs may also or in addition be characterized by the ability of two complementary nucleic acid strands to hybridize to each other under appropriately stringent conditions. Hybridizations are typically and preferably conducted with probe-length nucleic acid molecules, preferably 20-100 nucleotides in length. Those skilled in the art understand how to estimate and adjust the stringency of hybridization conditions such that sequences having at least a desired level of complementarity will stably hybridize, while those having lower complementarity will not. For examples of hybridization conditions and parameters, see, e.g., Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; Ausubel, F.M. et al. (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J. Homologs and homologous gene sequences may thus be identified using any nucleic acid sequence of interest, including the phage ORFs and bacterial target genes of the present invention.

A typical hybridization, for example, utilizes, besides the labeled probe of interest, a salt solution such as 6xSSC (NaCl and Sodium Citrate base) to stabilize nucleic acid strand interaction, a mild detergent such as 0.5% SDS, together with other typical additives such as Denhardt's solution and salmon sperm DNA. The solution is added to the immobilized sequence to be probed and incubated at suitable temperatures to preferably permit specific binding while minimizing nonspecific binding. The temperature of the incubations and ensuing washes is critical to the success and clarity of the hybridization. Stringent conditions employ relatively higher temperatures, lower salt concentrations, and/or more detergent than do non-stringent conditions. Hybridization temperatures also depend on the length, complementarity level, and nature (ie, "GC content") of the sequences to be tested. Typical stringent hybridizations and washes are conducted at temperatures of at least 40°C, while lower stringency hybridizations and washes are typically conducted at 37°C down to room temperature (~25°C). One of skill in the art is aware that these conditions may vary according to the parameters indicated above, and that certain additives such as formamide and dextran sulphate may also be added to affect the conditions.

By "stringent hybridization conditions" is meant hybridization conditions at least as stringent as the following: hybridization in 50% formamide, 5X SSC, 50 mM NaH₂PO₄, pH 6.8, 0.5% SDS, 0.1 mg/mL sonicated salmon sperm DNA, and 5X Denhart's solution at 42°C overnight; washing with 2X SSC, 0.1% SDS at 45°C; and washing with 0.2X SSC, 0.1% SDS at 45°C.

In sequence comparison analyses, an ORF, or motif, or set of motifs in a bacteriophage sequence can be compared to known inhibitor sequences, *e.g.*, homologous sequences encoding homologous inhibitors of bacterial function. Likewise, the analysis can include comparison with the structure of essential bacterial gene products, as structural similarities can be indicative of similar or replacement biological function. Such analysis can include the identification of a signature, or characteristic motif(s) of an inhibitor or inhibitor class.

Also, the identification of structural motifs in an encoded product, based on nucleotide or amino acid sequence analysis, can be used to infer a biochemical function for the product. A database containing identified structural motifs in a large number of sequences is available for identification of motifs in phage sequences. The database is PROSITE, which is available at www.expasy.ch/cgi-bin/scanprosite. The identification of motifs can, for example, include the identification of signature motifs for a class or classes of inhibitory proteins. Other such databases may also be used.

In aspects and preferred embodiments described herein, in which a bacterium or host bacterium is specified, the bacterium or host bacterium is preferably selected from a pathogenic bacterial species, for example, one selected from Table 1. Preferably, an animal or plant pathogen is used. For animals, preferably the bacterium is a bird or mammalian pathogen, still more preferably a human pathogen.

In aspects and preferred embodiments involving a bacteriophage or sequences from a bacteriophage, one or more bacteriophage are preferably selected from those listed in Table 1. Those exemplary bacteriophage are readily obtained from the indicated sources.

In some cases, it is advantageous to utilize phage with non-pathogenic host bacteria. The genome, structural motif, ORF, homolog, and other analyses described herein can be performed on such phage and bacteria. Such analysis provides useful information and compositions. The results of such analyses can also be utilized in aspects of the present invention to identify homologous ORFs, especially inhibitor ORFs in phage with pathogenic bacterial hosts. Similarly, identification of a target in a non-pathogenic host can be used to identify homologous sequences and targets in pathogenic bacteria, especially in genetically closely related bacteria. Those skilled in the art are familiar with bacterial genetic relationships and with how to determine relatedness based on levels of genomic identity or other measures of nucleotide sequence and/or amino acid sequence similarity, and/or other physical and culture characteristics such as morphology, nutritional requirements, or minimal media to support growth.

Also in preferred embodiments, an embodiment of this aspect is combined with an embodiment of the following aspect.

A related aspect of the invention provides methods for identifying a target for antibacterial agents by identifying the bacterial target(s) of at least one
5 uncharacterized or untargeted inhibitor protein or RNA from a bacteriophage. Such identification allows the development of antibacterial agents active on such targets. Preferred embodiments for identifying such targets involve the identification of binding of target and phage ORF products to one another. The phage ORF products may be subportions of a larger ORF product that also binds the host target. In
10 preferred embodiments, the phage protein or RNA is from an uncharacterized bacteriophage in Table 1. This aspect preferably includes the identification of a plurality of such targets in one or a plurality of different bacteria, preferably in one or a plurality of bacteria listed in Table 1.

In preferred embodiments of this aspect and other aspects of this invention
15 involving particular phage ORFs or phage sequences, the ORF is *Staphylococcus aureus* phage 77 ORF 17, 19, 43, 102, 104, or 182 as identified in U.S. application 09/407,804, *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

20 As indicated for the above aspect, preferably the method involves the use of a plurality of different phage, and thus a plurality of different phage inhibitors and/or inhibitor ORFs.

In addition to uncharacterized phage ORF products, it is also useful to identify the targets of phage ORF products which are known to be inhibitors of host bacteria,
25 but where the target has not been identified. Thus, such inhibitors can likewise be utilized as "untargeted" inhibitor phage ORFs and ORF products, e.g., proteins or RNAs.

In the context of inhibitor proteins or RNAs from a phage, the term "uncharacterized" means that a bacteria-inhibiting function for the protein has not
30 previously been identified. Preferably, but not necessarily, the sequence of the protein or the corresponding coding region or ORF was not described in the art before the filing of the present application for patent (or alternatively prior to the present invention). Thus, this term specifically excludes any bacteria-inhibiting phage protein and its associated bacterial target which has been identified as inhibitory before the
35 present invention or alternatively before the filing of the present application, for example those identified in Tables 12-14 or otherwise identified herein. For example, from *E. coli*, phage T7 genes 0.7 and 2.0 target the host RNA polymerase, phage T4

gp55/gp33 alter the specificity of host RNA polymerase. The T4 *regB* gene product also targets the host translation apparatus. As with the uncharacterized bacteriophage ORFs or bacteriophage above, for such identified proteins, the sequences encoding those proteins are excluded from the uncharacterized inhibitor proteins.

5 The term "fragment" refers to a portion of a larger molecule or assembly. For proteins, the term "fragment" refers to a molecule which includes at least 5 contiguous amino acids from the reference polypeptide or protein, preferably at least 8, 10, 12, 15, 20, 30, 50 or more contiguous amino acids. In connection with oligo- or polynucleotides, the term "fragment" refers to a molecule which includes at least 15
10 contiguous nucleotides from a reference polynucleotide, preferably at least 24, 30, 36, 45, 60, 90, 150, or more contiguous nucleotides.

 Preferred embodiments involve identification of binding that include methods for distinguishing bound molecules, for example, affinity chromatography, immunoprecipitation, crosslinking, and/or genetic screen methods that permit
15 protein:protein interactions to be monitored. One of skill in the art is familiar with these techniques and common materials utilized (see, *e.g.*, Coligan, J. et al. (eds.) (1995) Current Protocols in Protein Science, John Wiley & Sons, Secaucus, N.J.).

 Genetic screening for the identification of protein:protein interactions typically involves the co-introduction of both a chimeric bait nucleic acid sequence (here, the
20 phage ORF to be tested) and a chimeric target nucleic acid sequence that, when co-expressed and having affinity for one another in a host cell, stimulate reporter gene expression to indicate the relationship. A "positive" can thus suggest a potential inhibitory effect in bacteria. This is discussed in further detail in the Detailed Description section below. In this way, new bacterial targets can be identified that are
25 inhibited by specific phage ORF products or derivatives, fragments, mimetics, or other molecules.

 Other embodiments involve the identification and/or utilization of mutant targets by virtue of their host's relatively unresponsive nature in the presence of expression of ORFs previously identified as inhibitory to the non-mutant or wild-type
30 strain. Such mutants have the effect of protecting the host from an inhibition that would otherwise occur and indirectly allow identification of the precise responsible target for follow-up studies and anti-microbial development. In certain embodiments, rescue from inhibition occurs under conditions in which a bacterial target or mutant target is highly expressed. This is performed, for example, through coupling of the
35 sequence with regulatory element promoters, *e.g.*, as known in the art, which regulate expression at levels higher than wild-type, *e.g.*, at a level sufficiently higher than the

inhibitor can be competitively bound to the highly expressed target such that the bacterium is detectably less inhibited.

Identification of the bacterial target can involve identification of a phage-specific site of action. This can involve a newly identified target, or a target where the phage site of action differs from the site of action of a previously known antibacterial agent or inhibitor. For example, phage T7 genes 0.7 and 2.0 target the host RNA polymerase, which is also the cellular target for the antibacterial agent, rifampin. To the extent that a phage product is found to act at a different site than previously described inhibitors, aspects of the present invention can utilize those new, phage-specific sites for identification and use of new agents. The site of action can be identified by techniques well-known to those skilled in the art, for example, by mutational analysis, binding competition analysis, and/or other appropriate techniques.

Once a bacterial host target protein or nucleic acid or mutant target sequence has been identified and/or isolated, it too can be conveniently sequenced, sequence analyzed (*e.g.*, by computer), and the underlying gene(s), and corresponding translated product(s) further characterized. Preferred embodiments include such analysis and identification. Preferably such a target has not previously been identified as an appropriate target for antibacterial action.

Certain embodiments include the identification of at least one inhibitory phage ORF or ORF product, *e.g.*, as described for the above aspect, and thus are a combination of the two aspects.

Additionally, the invention provides methods for identifying targets for antibacterial agents by identifying homologs of a bacterial target *e.g.*, *S. aureus*, *Enterococcus faecalis* or other *Enterococci*, and *Streptococcus pneumoniae* of a bacteriophage inhibitory ORF product. Such homologs may be utilized in the various aspects and embodiments described herein as described for the host *Enterococcus* sp. for bacteriophage 182.

Other aspects of the invention provide isolated, purified, or enriched specific phage nucleic acid and amino acid sequences, subsequences, and homologs thereof for phage selected from uncharacterized phage listed in Table 1, preferably from bacteriophage 77, 3A, 96, 44AHJD (*Staphylococcus aureus* host bacterium), Dp-1 (*Streptococcus pneumoniae* host), or 182 (*Enterococcus* host) or other phage listed in Table 1 for those bacteria. For example, such sequences do not include sequences identified in any of Tables 11-14. Nucleotide sequences of this aspect are at least 15 nucleotides in length, preferably at least 18, 21, 24, or 27 nucleotides in length, more preferably at least 30, 50, or 90 nucleotides in length. In certain embodiments, longer

nucleic acids are preferred, for example those of at least 120, 150, 200, 300, 600, 900 or more nucleotides. Such sequences can, for example, be amplification oligonucleotides (*e.g.*, PCR primers), oligonucleotide probes, sequences encoding a portion or all of a phage-encoded protein, or a fragment or all of a phage-encoded protein. In preferred embodiments, the nucleic acid sequence contains a sequence which is within a length range with a lower length as specified above, and an upper length limit which is no more than 50, 60, 70, 80, or 90% of the length of the corresponding full-length ORF. The upper length limit can also be expressed in terms of the number of base pairs of the ORF (coding region). In preferred embodiments, the nucleic acid sequence is from *Staphylococcus aureus* phage 77 ORF 17, 19, 43, 102, 104, or 182 as identified in U.S. application 09/407,804, *S. aureus* phage 44 AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

As it is recognized that alternate codons will encode the same amino acid for most amino acids due to the degeneracy of the genetic code, the sequences of this aspect includes nucleic acid sequences utilizing such alternate codon usage for one or more codons of a coding sequence. For example, all four nucleic acid sequences GCT, GCC, GCA, and GCG encode the amino acid, alanine. Therefore, if for an amino acid there exists an average of three codons, a polypeptide of 100 amino acids in length will, on average, be encoded by 3^{100} , or 5×10^{47} , nucleic acid sequences. Thus, a nucleic acid sequence can be modified (*e.g.*, a nucleic acid sequence from a phage as specified above) to form a second nucleic acid sequence encoding the same polypeptide as encoded by the first nucleic acid sequence using routine procedures and without undue experimentation. Thus, all possible nucleic acid sequences that encode the specified amino acid sequences are also fully described herein, as if all were written out in full, taking into account the codon usage, especially that preferred in the host bacterium. The alternate codon descriptions are available in common textbooks, for example, Stryer, BIOCHEMISTRY 3rd ed., and Lehninger, BIOCHEMISTRY 3rd ed., along with many others. Codon preference tables for various types of organisms are available in the literature. Sequences with alternate codons at one or more sites can also be utilized in the computer-related aspects and embodiments herein. Because of the number of sequence variations involving alternate codon usage, for the sake of brevity, individual sequences are not separately listed herein. Instead the alternate sequences are described by reference to the natural sequence with replacement of one or more (up to all *e.g.*, up to 3, 5, 10, 15, 20, 30, 40, 50, or more) of the degenerate codons with alternate codons from the alternate codon

table (Table 6), or a modified table applicable to a particular organism that has differing codon usage, preferably with selection according to preferred codon usage for the normal host organism or a host organism in which a sequence is intended to be expressed. Those skilled in the art also understand how to alter the alternate codons to be used for expression in organisms where certain codons code differently than shown in the "universal" codon table.

For amino acid sequences or polypeptides, sequences contain at least 5 peptide-linked amino acid residues, and preferably at least 6, 7, 10, 15, 20, 30, or 40, amino acids having identical amino acid sequence as the same number of contiguous amino acid residues in a particular phage ORF product. In some cases longer sequences may be preferred, for example, those of at least 50, 60, 70, 80, or 100 amino acids in length. In preferred embodiments, the amino acid sequence contains a sequence which is within a length range with a lower length as specified above, and an upper length limit which is no more than 50, 60, 70, 80, or 90% of the length of the corresponding full-length ORF product. The upper length limit can also be expressed in terms of the number of amino acid residues of the ORF product. In preferred embodiments, the amino acid sequence or polypeptide has bacteria-inhibiting function when expressed or otherwise present in a bacterial cell which is a host for the bacteriophage from which the sequence was derived.

By "isolated" in reference to a nucleic acid is meant that a naturally occurring sequence has been removed from its normal cellular (*e.g.*, chromosomal) environment or is synthesized in a non-natural environment (*e.g.*, artificially synthesized). Thus, the sequence may be in a cell-free solution or placed in a different cellular environment. The term does not imply that the sequence is the only nucleotide chain present, but that it is essentially free (about 90-95% pure at least) of non-nucleotide material naturally associated with it, and thus is distinguished from isolated chromosomes.

The term "enriched" means that the specific DNA or RNA sequence constitutes a significantly higher fraction (2-5 fold) of the total DNA or RNA present in the cells or solution of interest than in normal or diseased cells or in cells from which the sequence was originally taken. This could be caused by a person by preferential reduction in the amount of other DNA or RNA present, or by a preferential increase in the amount of the specific DNA or RNA sequence, or by a combination of the two. However, it should be noted that enriched does not imply that there are no other DNA or RNA sequences present, just that the relative amount of the sequence of interest has been significantly increased.

The term "significant" is used to indicate that the level of increase is useful to the person making such an increase and an increase relative to other nucleic acids of about at least 2-fold, more preferably at least 5- to 10-fold or even more. The term also does not imply that there is no DNA or RNA from other sources. The other
5 source DNA may, for example, comprise DNA from a yeast or bacterial genome, or a cloning vector such as pUC19. This term distinguishes from naturally occurring events, such as viral infection, or tumor type growths, in which the level of one mRNA may be naturally increased relative to other species of mRNA. That is, the term is meant to cover only those situations in which a person has intervened to
10 elevate the proportion of the desired nucleic acid.

It is also advantageous for some purposes that a nucleotide sequence be in purified form. The term "purified" in reference to nucleic acid does not require absolute purity (such as a homogeneous preparation). Instead, it represents an indication that the sequence is relatively more pure than in the natural environment
15 (compared to the natural level, this level should be at least 2-5 fold greater, *e.g.*, in terms of mg/mL). Individual clones isolated from a cDNA library may be purified to electrophoretic homogeneity. The claimed DNA molecules obtained from these clones could be obtained directly from total DNA or from total RNA. The cDNA clones are not naturally occurring, but rather are preferably obtained via manipulation
20 of a partially purified naturally occurring substance (messenger RNA). The construction of a cDNA library from mRNA involves the creation of a synthetic substance (cDNA) and pure individual cDNA clones can be isolated from the synthetic library by clonal selection of the cells carrying the cDNA library. Thus, the process which includes the construction of a cDNA library from mRNA and isolation
25 of distinct cDNA clones yields an approximately 10^6 -fold purification of the native message. Thus, purification of at least one order of magnitude, preferably two or three orders, and more preferably four or five orders of magnitude is expressly contemplated.

The terms "isolated", "enriched", and "purified" as respect nucleic acids,
30 above, may similarly be used to denote the relative purity and abundance of polypeptides (multimers of amino acids joined one to another by α -carboxyl: α -amino group (peptide) bonds). These, too, may be stored in, grown in, screened in, and selected from libraries using biochemical techniques familiar in the art. Such polypeptides may be natural, synthetic or chimeric and may be extracted using any of
35 a variety of methods, such as antibody immunoprecipitation, other "tagging" techniques, conventional chromatography and/or electrophoretic methods. Some of the above utilize the corresponding nucleic acid sequence.

As indicated above, aspects and embodiments of the invention are not limited to entire genes and proteins. The invention also provides and utilizes fragments and portions thereof, preferably those which are "active" in the inhibitory sense described above. Such peptides or oligopeptides and oligo or polynucleotides have preferred
5 lengths as specified above for nucleic acid and amino acid sequences from phage; corresponding recombinant constructs can be made to express the encoded same. Also included are homologous sequences and fragments thereof.

Nucleic acid sequences of the present invention can be isolated using a method similar to those described herein or other methods known to those skilled in the art.
10 In addition, such nucleic acid sequences can be chemically synthesized by well-known methods. Also, by having particular phage ORFs, e.g., the phage ORFs identified herein (e.g., anti-bacterial ORFs of the present invention, portions thereof, or oligonucleotides derived therefrom as described), other antimicrobial sequences from other bacteriophage sources can be identified and isolated using methods
15 described here or other methods, including methods utilizing nucleic acid hybridization and/or computer-based sequence alignment methods.

The invention also provides bacteriophage antimicrobial DNA segments from other phages based on nucleic acids and sequences hybridizing to the presently identified inhibitory ORF under high stringency conditions or sequences that are
20 highly homologous. The bacteriophage segment from a specific phage, e.g., an antimicrobial DNA segment, can be used to identify a related segment from another unrelated phage based on stringent conditions of hybridization or on being a homolog based on nucleic acid and/or amino acid sequence comparisons. As with identified inhibitory sequences, such homologous coding sequences and products can be used as
25 antimicrobials, to construct active portions or derivatives, to construct peptidomimetics, and to identify bacterial targets.

The nucleotide and amino acid sequences identified herein are believed to be correct, however, certain sequences may contain a small percentage of errors, e.g., 1-5%. In the event that any of the sequences have errors, the corrected sequences can be
30 readily provided by one skilled in the art using routine methods. For example, the nucleotide sequences can be confirmed or corrected by obtaining and culturing the relevant phage, and purifying phage genomic nucleic acids. A region or regions of interest can be amplified, e.g., by PCR from the appropriate genomic template, using primers based on the described sequence. The amplified regions can then be
35 sequenced using any of the available methods (e.g., a dideoxy termination method).

This can be done redundantly to provide the corrected sequence or to confirm that the described sequence is correct. Alternatively, a particular sequence or sequences can be identified and isolated as an insert or inserts in a phage genomic library and isolated, amplified, and sequenced by standard methods. Confirmation or correction of a nucleotide sequence for a phage gene provides an amino acid sequence of the encoded product by merely reading off the amino acid sequence according to the normal codon relationships and/or expressed in a standard expression system and the polypeptide product sequenced by standard techniques. The sequences described herein thus provide unique identification of the corresponding genes, coding sequences, and other sequences, allowing those sequences to be used in the various aspects of the present invention.

In other aspects, the invention provides recombinant vectors and cells harboring at least one of the phage ORFs or portion thereof, or bacterial target sequences described herein. As understood by those skilled in the art, vectors may be provided in different forms, including, for example, plasmids, cosmids, and virus-based vectors. See, *e.g.*, Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; See also, Ausubel, F.M. et al. (eds.) (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J.

In preferred embodiments, the vectors will be expression vectors, preferably shuttle vectors that permit cloning, replication, and expression within bacteria. An "expression vector" is one having regulatory nucleotide sequences containing transcriptional and translational regulatory information that controls expression of the nucleotide sequence in a host cell. Preferably the vector is constructed to allow amplification from vector sequences flanking an insert locus. In certain embodiments, the expression vectors may additionally or alternatively support expression, and/or replication in animal, plant and/or yeast cells due to the presence of suitable regulatory sequences, *e.g.*, promoters, enhancers, 3' stabilizing sequences, primer sequences, etc. In preferred embodiments, the promoters are inducible and specific for the system in which expression is desired, *e.g.*, bacteria, animal, plant, or yeast. The vectors may optionally encode a "tag" sequence or sequences to facilitate protein purification. Convenient restriction enzyme cloning sites and suitable selective marker(s) are also optionally included. Such selective markers can be, for example, antibiotic resistance markers or markers which supply an essential nutritive growth factor to an otherwise deficient mutant host, *e.g.*, tryptophan, histidine, or leucine in the Yeast Two-Hybrid systems described below.

The term "recombinant vector" relates to a single- or double-stranded circular nucleic acid molecule that can be transfected into cells and replicated within or independently of a cell genome. A circular double-stranded nucleic acid molecule can be cut and thereby linearized upon treatment with appropriate restriction enzymes. An assortment of nucleic acid vectors, restriction enzymes, and the knowledge of the nucleotide sequences cut by restriction enzymes are readily available to those skilled in the art. A nucleic acid molecule encoding a desired product can be inserted into a vector by cutting the vector with restriction enzymes and ligating the two pieces together. Preferably the vector is an expression vector, *e.g.*, a shuttle expression vector as described above.

By "recombinant cell" is meant a cell possessing introduced or engineered nucleic acid sequences, *e.g.*, as described above. The sequence may be in the form of or part of a vector or may be integrated into the host cell genome. Preferably the cell is a bacterial cell.

In another aspect, the invention also provides methods for identifying and/or screening compounds "active on" at least one bacterial target of a bacteriophage inhibitor protein or RNA. Preferred embodiments involve contacting such a bacterial target or targets (*e.g.*, bacterial target proteins) with a test compound, and determining whether the compound binds to or reduces the level of activity of the bacterial target (*e.g.*, a bacterial target protein). Preferably this is done either *in vivo* (*i.e.*, in a cell-based assay) or *in vitro*, *e.g.*, in a cell-free system under approximately physiological conditions.

The compounds that can be used may be large or small, synthetic or natural, organic or inorganic, proteinaceous or non-proteinaceous. In preferred embodiments, the compound is a peptidomimetic, as described herein, a bacteriophage inhibitor protein or fragment or derivative thereof, preferably an "active portion", or a small molecule.

In preferred embodiments, the bacterial target is a target of a phage ORF identified herein, *e.g.*, *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

In particular embodiments, the methods include the identification of bacterial targets or the site of action of an inhibitor on a bacterial target as described above or otherwise described herein.

In embodiments involving binding assays, preferably binding is to a fragment or portion of a bacterial target protein, where the fragment includes less than 90%, 80%, 70%, 60%, 50%, 40%, or 30% of an intact bacterial target protein. Preferably,

the at least one bacterial target includes a plurality of different targets of bacteriophage inhibitor proteins, preferably a plurality of different targets. The plurality of targets can be in or from a plurality of different bacteria, but preferably is from a single bacterial species.

5 A "method of screening" refers to a method for evaluating a relevant activity or property of a large plurality of compounds (e.g., a bacteria-inhibiting activity), rather than just one or a few compounds. For example, a method of screening can be used to conveniently test at least 100, more preferably at least 1000, still more preferably at least 10,000, and most preferably at least 100,000 different compounds,
10 or even more.

In the context of this invention, the term "small molecule" refers to compounds having molecular mass of less than 2000 Daltons, preferably less than 1500, still more preferably less than 1000, and most preferably less than 600 Daltons. Preferably but not necessarily, a small molecule is not an oligopeptide.

15 In a related aspect or in preferred embodiments, the invention provides a method of screening for potential antibacterial agents by determining whether any of a plurality of compounds, preferably a plurality of small molecules, is active on at least one target of a bacteriophage inhibitor protein or RNA. Preferred embodiments include those described for the above aspect, including embodiments which involve
20 determining whether one or more test compounds bind to or reduce the level of activity of a bacterial target, and embodiments which utilize a plurality of different targets as described above.

The identification of bacteria-inhibiting phage ORFs and their encoded products also provides a method for identifying an active portion of such an encoded
25 product. This also provides a method for identifying a potential antibacterial agent by identifying such an active portion of a phage ORF or ORF product. In preferred embodiments, the identification of an active portion involves one or more of mutational analysis, deletion analysis, or analysis of fragments of such products. The method can also include determination of a 3-dimensional structure of an active
30 portion, such as by analysis of crystal diffraction patterns. In further embodiments, the method involves constructing or synthesizing a peptidomimetic compound, where the structure of the peptidomimetic compound corresponds to the structure of the active portion. In this context, "corresponds" means that the peptidomimetic compound structure has sufficient similarities to the structure of the active portion that
35 the peptidomimetic will interact with the same molecule as the phage protein and preferably will elicit at least one cellular response in common which relates to the inhibition of the cell by the phage protein.

In preferred embodiments, the ORF or ORF product is or is derived or obtained from *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014 or product thereof.

5 The methods for identifying or screening for compounds or agents active on a bacterial target of a phage-encoded inhibitor can also involve identification of a phage-specific site of action on the target.

Preferably in the methods for identifying or screening for compounds active on such a bacterial target, the target is uncharacterized; the target is from an uncharacterized bacterium from Table 1; the site of action is a phage-specific site of action.

Further embodiments include the identification of inhibitor phage ORFs and bacterial targets as in aspects above.

15 An "active portion" as used herein denotes an epitope, a catalytic or regulatory domain, or a fragment of a bacteriophage inhibitor protein that is responsible for, or a significant factor in, bacterial target inhibition. The active portion preferably may be removed from its contiguous sequences and, in isolation, still effect inhibition.

By "mimetic" is meant a compound structurally and functionally related to a reference compound that can be natural, synthetic, or chimeric. In terms of the present invention, a "peptidomimetic," for example, is a compound that mimics the activity-related aspects of the 3-dimensional structure of a peptide or polypeptide in a non-peptide compound, for example mimics the structure of a peptide or active portion of a phage- or bacterial ORF-encoded polypeptide.

25 A related aspect provides a method for inhibiting a bacterial cell by contacting the bacterial cell with a compound active on a bacterial target of a bacteriophage inhibitor protein or RNA, where the target was uncharacterized. In preferred embodiments, the compound is such a protein, or a fragment or derivative thereof; a structural mimetic, *e.g.*, a peptidomimetic, of such a protein or fragment; a small molecule; the contacting is performed *in vitro*, the contacting is performed *in vivo* in an infected or at risk organism, *e.g.*, an animal such as a mammal or bird, for example, a human, or other mammal described herein; the bacterium is selected from a genus and/or species listed in Table 1; the bacteriophage inhibitor protein is uncharacterized; the bacteriophage inhibitor protein is from an uncharacterized phage listed in Table 1; the phage inhibitor protein is from one of *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

In the context of targets in this invention, the term "uncharacterized" means that the target was not recognized as an appropriate target for an antibacterial agent prior to the filing of the present application or alternatively prior to the present invention. Such lack of recognition can include, for example, situations where the target and/or a nucleotide sequence encoding the target were unknown, situations where the target was known, but where it had not been identified as an appropriate target or as an essential cellular component, and situations where the target was known as essential but had not been recognized as an appropriate target due to a belief that the target would be inaccessible or otherwise that contacting the cell with a compound active on the target *in vitro* would be ineffective in cellular inhibition, or ineffective in treatment of an infection. Methods described herein utilizing bacterial targets, *e.g.*, for inhibiting bacteria or treating bacterial infections, can also utilize "uncharacterized target sites", meaning that the target has been previously recognized as an appropriate target for an antibacterial agent, but where an agent or inhibitor of the invention is used which acts at a different site than that at which the previously utilized antibacterial agent, *i.e.*, a phage-specific site. Preferably the phage-specific site has different functional characteristics from the previously utilized site. In the context of targets or target sites, the term "phage-specific" indicates that the target or site is utilized by at least one bacteriophage as an inhibitory target and is different from previously identified targets or target sites.

In the context of this invention, the term "bacteriophage inhibitor protein" refers to a protein encoded by a bacteriophage nucleic acid sequence which inhibits bacterial function in a host bacterium. Thus, it is a bacteria-inhibiting phage product.

In the context of this invention, the phrase "contacting the bacterial cell with a compound active on a bacterial target of a bacteriophage inhibitor protein" or equivalent phrases refer to contacting with an isolated, purified, or enriched compound or a composition including such a compound, but specifically does not rely on contacting the bacterial cell with an intact phage which encodes the compound. Preferably no intact phage are involved in the contacting.

Related aspects provide methods for prophylactic or therapeutic treatment of a bacterial infection by administering to an infected, challenged or at risk organism a therapeutically or prophylactically effective amount of a compound active on a target of a bacteriophage inhibitor protein or RNA, or as described for the previous aspect. Preferably the bacterium involved in the infection or risk of infection produces the identified target of the bacteriophage inhibitor protein or alternatively produces a homologous target compound. In preferred embodiments, the host organism is a plant or animal, preferably a mammal or bird, and more preferably, a human or other

mammal described herein. Preferred embodiments include, without limitation, those as described for the preceding aspect.

Compounds useful for the methods of inhibiting, methods of treating, and pharmaceutical compositions can include novel compounds, but can also include
5 compounds which had previously been identified for a purpose other than inhibition of bacteria. Such compounds can be utilized as described and can be included in pharmaceutical compositions.

In preferred embodiments of this and other aspects of the invention utilizing bacterial target sequences of a bacteriophage inhibitory ORF product, the target
10 sequence is encoded by a *Staphylococcus* nucleic acid coding sequence, preferably *S. aureus*, a *Streptococcus* nucleic acid coding sequence, preferably *Streptococcus pneumoniae*, or *Enterococcus* nucleic acid coding sequence. Possible target sequences are described herein by reference to sequence source sites.

The amino acid sequence of a polypeptide target is readily provided by
15 translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. For the sake of brevity, the sequences are described by reference to the GenBank entries instead of being written out in full herein. In cases where the TIGR or GenBank entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a
20 phage host genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

25 In the context of nucleic acid or amino acid sequences of this invention, the term "corresponding" indicates that the sequence is at least 95% identical, preferably at least 97% identical, and more preferably at least 99% identical to a sequence from the specified phage genome, a ribonucleotide equivalent, a degenerate equivalent (utilizing one or more degenerate codons), or a homologous sequence, where the
30 homolog provides functionally equivalent biological function.

By "treatment" or "treating" is meant administering a compound or pharmaceutical composition for prophylactic and/or therapeutic purposes. The term "prophylactic treatment" refers to treating a patient or animal that is not yet infected but is susceptible to or otherwise at risk of a bacterial infection. The term "therapeutic
35 treatment" refers to administering treatment to a patient already suffering from infection.

The term "bacterial infection" refers to the invasion of the host organism, animal or plant, by pathogenic bacteria. This includes the excessive growth of bacteria which are normally present in or on the body of the organism, but more generally, a bacterial infection can be any situation in which the presence of a bacterial population(s) is damaging to a host organism. Thus, for example, an organism suffers from a bacterial population when excessive numbers of a bacterial population are present in or on the organism's body, or when the effects of the presence of a bacterial population(s) is damaging to the cells, tissue, or organs of the organism.

The terms "administer", "administering", and "administration" refer to a method of giving a dosage of a compound or composition, *e.g.*, an antibacterial pharmaceutical composition, to an organism. Where the organism is a mammal, the method is, *e.g.*, topical, oral, intravenous, transdermal, intraperitoneal, intramuscular, or intrathecal. The preferred method of administration can vary depending on various factors, *e.g.*, the components of the pharmaceutical composition, the site of the potential or actual bacterial infection, the bacterium involved, and the infection severity.

The term "mammal" has its usual biological meaning referring to any organism of the Class Mammalia of higher vertebrates that nourish their young with milk secreted by mammary glands, *e.g.*, mouse, rat, and, in particular, human, bovine, sheep, swine, dog, and cat.

In the context of treating a bacterial infection a "therapeutically effective amount" or "pharmaceutically effective amount" indicates an amount of an antibacterial agent, *e.g.*, as disclosed for this invention, which has a therapeutic effect. This generally refers to the inhibition, to some extent, of the normal cellular functioning of bacterial cells that renders or contributes to bacterial infection.

The dose of antibacterial agent that is useful as a treatment is a "therapeutically effective amount." Thus, as used herein, a therapeutically effective amount means an amount of an antibacterial agent that produces the desired therapeutic effect as judged by clinical trial results and/or animal models. This amount can be routinely determined by one skilled in the art and will vary depending on several factors, such as the particular bacterial strain involved and the particular antibacterial agent used.

In connection with claims to methods of inhibiting bacteria and therapeutic or prophylactic treatments, "a compound active on a target of a bacteriophage inhibitor protein" or terms of equivalent meaning differ from administration of or contact with an intact phage naturally encoding the full-length inhibitor compound. While an intact phage may conceivably be incorporated in the present methods, the method at

least includes the use of an active compound as specified different from a full length inhibitor protein naturally encoded by a bacteriophage and/or a delivery or contacting method different from administration of or contact with an intact phage encoding the full-length protein. Similarly, pharmaceutical compositions described herein at least
5 include an active compound different from a full-length inhibitor protein naturally encoded by a bacteriophage or such a full-length protein is provided in the composition in a form different from being encoded by an intact phage. Preferably the methods and compositions do not include an intact phage.

In accord with the above aspects, the invention also provides antibacterial
10 agents and compounds active on bacterial targets of bacteriophage inhibitor proteins or RNAs, where the target was uncharacterized as indicated above. As previously indicated, such active compounds include both novel compounds and compounds which had previously been identified for a purpose other than inhibition of bacteria. Such previously identified biologically active compounds can be used in
15 embodiments of the above methods of inhibiting and treating. In preferred embodiments, the targets, bacteriophage, and active compound are as described herein for methods of inhibiting and methods of treating. Preferably the agent or compound is formulated in a pharmaceutical composition which includes a pharmaceutically acceptable carrier, excipient, or diluent. In addition, the invention provides agents,
20 compounds, and pharmaceutical compositions where an active compound is active on an uncharacterized phage-specific site.

In preferred embodiments, the target is as described for embodiments of aspects above.

Likewise, the invention provides a method of making an antibacterial agent.
25 The method involves identifying a target of a bacteriophage inhibitor polypeptide or protein or RNA, screening a plurality of compounds to identify a compound active on the target, and synthesizing the compound in an amount sufficient to provide a therapeutic effect when administered to an organism infected by a bacterium naturally producing the target. In preferred embodiments, the identification of the target and
30 identification of active compounds include steps or methods and/or components as described above (or otherwise herein) for such identification. Likewise, the active compound can be as described above, including fragments and derivatives of phage inhibitor proteins, peptidomimetics, and small molecules. As recognized by those skilled in the art, peptides can be synthesized by expression systems and purified, or
35 can be synthesized artificially. In preferred embodiments the inhibitory phage ORF products is from *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus*

pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

As indicated above, sequence analysis of nucleotide and/or amino acid sequences can beneficially utilize computer analysis. Thus, in additional aspects the invention provides computer-related hardware and media and methods utilizing and incorporating sequence data from uncharacterized phage, *e.g.*, uncharacterized phage listed in Table 1, preferably at least one of *Staphylococcus aureus* phage *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014, or 44 AHJD, *Enterococcus* sp. phage 182, or *Streptococcus pneumoniae* phage Dp-1. In general, such aspects can facilitate the above-described aspects. Various embodiments involve the analysis of genetic sequence and encoded products, as applied to the evaluating bacteriophage inhibitor ORFs and compounds and fragments related thereto. The various sequence analyses, as well as function analyses, can be used separately or in combination, as well as in preceding aspects and embodiments. Use in combination is often advantageous as the additional information allows more efficient prioritizing of phage ORFs for identification of those ORFs that provide bacteria-inhibiting function.

In one aspect, the invention provides a computer-readable device which includes at least one recorded amino acid or nucleotide sequence corresponding to one of the specified phage and a sequence analysis program for analyzing a nucleotide and/or amino acid sequence. The device is arranged such that the sequence information can be retrieved and analyzed using the analysis program. The analysis can identify, for example, homologous sequences or the indicated %s of the phage genome and structural motifs. Preferably the sequence includes at least 1 phage ORF or encoded product, more preferably at least 10%, 20%, 30%, 40%, 50%, 70%, 90%, or 100% of the genomic phage ORFs and/or equivalent cDNA, RNA, or amino acid sequences. Preferably the sequence or sequences in the device are recorded in a medium such as a floppy disk, a computer hard drive, an optical disk, computer random access memory (RAM), or magnetic tape. The program may also be recorded in such medium. The sequences can also include sequences from a plurality of different phage.

In this context, the term "corresponding" indicates that the sequence is at least 95% identical, preferably at least 97% identical, and more preferably at least 99% identical to a sequence from the specified phage genome, a ribonucleotide equivalent, a degenerate equivalent (utilizing one or more degenerate codons), or a homologous sequence, where the homolog provides functionally equivalent biological function.

Similarly, the invention provides a computer analysis system for identifying biologically important portions of a bacteriophage genome. The system includes a data storage medium, *e.g.*, as identified above, which has recorded thereon a nucleotide sequence corresponding to at least a portion of at least one uncharacterized bacteriophage genome, a set of program instructions to allow searching of the sequence or sequences to analyze the sequence, and an output device where the portion includes at least the sequence length as specified in the preceding aspect. The output device is preferably a printer, a video display, or a recording medium. More than one output device may be included. For each of the present computer-related aspects, the bacteriophage are preferably selected from the uncharacterized phage listed in Table 1, more preferably from bacteriophage 77, 3A, 96, 44 AHJD (*S. aureus*), Dp-1 (*Streptococcus pneumoniae*), or 182 (*Enterococcus*).

In keeping with the computer device aspects, the invention also provides a method for identifying or characterizing a bacteriophage ORF by providing a computer-based system for analyzing nucleotide or amino acid sequences, *e.g.*, as describe above. The system includes a data storage medium which has recorded a sequences or sequences as described for the above devices, a set of instructions as in the preceding aspect, and an output device as in the preceding aspect. The method further involves analyzing at least one sequence, and outputting the analysis results to at least one output device.

In preferred embodiments, the analysis identifies a sequence similarity or homology with a sequence or sequences selected from bacterial ORFs encoding products with related biological function; ORFs encoding known inhibitors; and essential bacterial ORFs. Preferably the analysis identifies a probable biological function based on identification of structural elements or characteristic or signature motifs of an encoded product or on sequence similarity or homology. Preferably the uncharacterized bacteriophage is from Table 1, more preferably at least one of bacteriophage 77, 3A, 96, 44 AHJD (*S. aureus*), Dp-1 (*Streptococcus pneumoniae*), or 182 (*Enterococcus*). In preferred embodiments, the method also involves determining at least a portion of the nucleotide sequence of at least one uncharacterized bacteriophage as indicated, and recording that sequence on data storage medium of the computer-based system. In preferred embodiments, the analysis identifies a sequence similarity of homology with a *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

As used in the claims to describe the various inventive aspects and embodiments, "comprising" means including, but not limited to, whatever follows the word "comprising". Thus, use of the term "comprising" indicates that the listed elements are required or mandatory, but that other elements are optional and may or may not be present. By "consisting of" is meant including, and limited to, whatever follows the phrase "consisting of". Thus, the phrase "consisting of" indicates that the listed elements are required or mandatory, and that no other elements may be present. By "consisting essentially of" is meant including any elements listed after the phrase, and limited to other elements that do not interfere with or contribute to the activity or action specified in the disclosure for the listed elements. Thus, the phrase "consisting essentially of" indicates that the listed elements are required or mandatory, but that other elements are optional and may or may not be present depending upon whether or not they affect the activity or action of the listed elements.

Further embodiments will be apparent from the following Detailed Description and from the claims.

BRIEF DESCRIPTION OF THE DRAWINGS

FIGURE 1A and 1B are flow schematics showing the manipulations used to convert pT0021, an arsenite inducible vector containing the luciferase gene, into pTHA or pTM, two *ars* inducible vectors. Vector pTHA contains BamH I, Sal I, and Hind III cloning sites and a downstream HA epitope tag. Vector pTM contains Bam HI and Hind III cloning sites and no HA epitope tag.

FIGURE 2 is a schematic representation of the cloning steps involved to place the DNA segments of any of ORFs 17/ 19/ 43/ 102/104/182 or other sequences into pTHA to assess inhibitory potential. For subcloning into pTM or pT0021, Individual ORFs were amplified by the PCR using oligonucleotides targeting the ATG and stop codons of the ORFs. Using this strategy, Bam HI and Hind III sites were positioned immediately upstream or downstream, respectively of the start and stop codons of each ORF. Following digestion with Bam HI and Hind III, the PCR fragments were subcloned into the same sites of pT0021 or pTM. Clones were verified by PCR and direct sequencing.

FIGURE 3 shows a schematic representation of the functional assays used to characterize the bactericidal and bacteriostatic potential of all predicted ORFs (>33 amino acids) encoded by bacteriophage 77. Fig. 3A) Functional assay on semi-solid support media. Fig. 3B) Functional assay in liquid culture.

FIGURE 4A, B, and C is a bar graph showing the results of a screen in liquid media to assess bacteriostatic or bactericidal activity of 93 predicted ORFs (>33 amino acids) encoded by bacteriophage 77. Growth inhibition assays were performed as detailed in the Detailed Description. The relative growth of *Staphylococcus aureus* transformants harboring a given bacteriophage 77 ORF (identified on the bottom of the graph), in the absence or presence of arsenite, is plotted relative to growth of a *Staphylococcus aureus* transformant containing ORF 5, a non-toxic bacteriophage 77 ORF (which is set at 100%). Each bar represents the average obtained from three *Staph A* transformants grown in duplicate. Bacteriophage 77 ORFs showing significant growth inhibition consist of ORFs 17, 19, 102, 104, and 182.

FIGURE 5 shows a block diagram of major components of a general purpose computer.

FIGURE 6 shows an ORF map for *Streptococcus pneumoniae* bacteriophage Dp-1 showing the ORF identifiers, genomic locations, and orientations of the 85 identified ORFs that were found to have ribosomal binding sites and thus are expected to be expressed.

FIGURE 7 shows a schematic representation of the arsenite-inducible expression system present in a shuttle vector designed to express individual *Streptococcus* bacteriophage Dp-1 ORFs in *Streptococcus*. Various modifications can be readily made to such a vector, or other vectors can be readily constructed to provide inducible expression of ORFs in a particular host bacterium using well-known techniques.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The invention may be more clearly understood from the following description.

5 The tables will first be briefly described.

Table 1 is a listing of a large number of available bacteriophage that can be readily obtained and used in the present invention.

Table 2 shows the complete nucleotide sequence of the genome of *Staphylococcus aureus* bacteriophage 77.

10 Table 3 shows a list of all the ORFs from Bacteriophage 77 that were screened in the functional assay to identify those with anti-microbial activity.

Table 4 shows the predicted nucleotide sequence, predicted amino acid sequence, and physiochemical parameters of ORF 17/ 19/ 43/ 102/ 104/ 182]. These include the primary amino acid sequence of the predicted protein, the average
15 molecular weight, amino acid composition, theoretical pI, hydrophobicity map, and predicted secondary structure map.

Table 5 shows homology search results. BLAST analysis was performed with ORFs 17/ 19/ 43/ 102/ 104/ 182 against NCBI non-redundant nucleotide and Swissprot databases. The results of this search indicate that: I) ORF 17 has no
20 significant homology to any gene in the NCBI non-NCBI non-redundant nucleotide database, II) ORF 19 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 59 of bacteriophage phi PVL, III) ORF 43 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 39 of phi PVL, IV) ORF 102 has
25 significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 38 of phi PVL, V) ORF 104 has no significant homology to any gene in the NCBI non-redundant nucleotide database, VI) ORF 182 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 39 of phi PVL.

30 Table 6 is a table from Alberts et al., MOLECULAR BIOLOGY OF THE CELL 3rd ed., showing the redundancy of the "universal" genetic code.

Table 7 shows the complete nucleotide sequence of *Staphylococcus aureus* bacteriophage 3A.

Table 8 is a listing of the ORFs identified in *Staphylococcus aureus* bacteriophage 3A.

Table 9 shows the complete nucleotide sequence of *Staphylococcus aureus* bacteriophage 96.

5 Table 10 is a listing of the ORFs identified in *Staphylococcus aureus* bacteriophage 96.

Table 11 is a listing of sequences deposited in the NCBI public database (GeneBank) for bacteriophage listed in Table 1.

10 Table 12 is a listing of phage which encode a known lysis function , including the identified lysis gene.

Table 13 is a listing of bacteriophage which encode holin genes, where holin genes encode proteins which form pores and eventually enable other enzymes to kill the host bacterium.

Table 14 is a listing of bacteriophage which encode kil genes.

15 Table 15 is a list of *Staphylococcus aureus* sequences identified by accession number which may include sequences from genes coding for target sequences for the phage 77-encoded antimicrobial proteins or peptides. The sequences were obtained by searching GenBank for listings.

20 Table 16 shows the nucleotide sequence of the genome of *Staphylococcus aureus* phage 44 AHJD.

Table 17 lists and shows the sequence position of the 73 ORFs predicted to be encoded by *Staphylococcus aureus* bacteriophage 44 AHJD that are greater than 33 amino acids.

25 Table 18 shows the ORF sequences and putative amino acid sequences for the *Staphylococcus aureus* bacteriophage 44AHJD ORFs greater than 33 amino acids.

Table 19 shows the similarities in sequence identified between predicted *Staphylococcus aureus* bacteriophage 44 AHJD ORFs and sequences present in public databases.

30 Table 20 shows the homology alignments between predicted *Staphylococcus aureus* bacteriophage 44AHJD ORFs and the corresponding protein sequences present in public sequence databases.

Table 21 shows the complete nucleotide sequence of the genome of *Enterococcus* bacteriophage 182.

35 Table 22 lists and shows the sequence position of the 80 ORFs identified in bacteriophage 182 and that are greater than 33 amino acids.

Table 23 shows the nucleotide and predicted amino acid sequence of all 80 ORFs identified in bacteriophage 182.

Table 24 shows the similarities identified to date in sequence between *Enterococcus* phage 182 ORFs greater than 33 amino acids and sequences present in public sequence databases.

Table 25 shows the predicted amino acid sequence as well as the predicted secondary structures map for two *Enterococcus* bacteriophage 182 ORFs.

Table 26 shows the homology alignments between predicted *Enterococcus* bacteriophage 182 ORFs and the corresponding protein sequences present in public sequence databases.

Table 27 list *Enterococcus* sequences listed in GenBank providing possible Enterococcal target sequences for inhibitory *Enterococcus* bacteriophage 182 ORFs and other compounds with antibacterial activity.

Table 28 shows the complete nucleotide sequence of the genome of *Streptococcus* bacteriophage Dp-1.

Table 29 lists and shows sequence position of the 273 ORFs identified in Pneumococcal bacteriophage Dp-1 that are greater than 33 amino acids, 85 of which are predicted to be expressed in Dp-1 as having a ribosomal binding site. That set of 85 ORFs is shown in the attached drawings.

Table 30 shows the nucleotide and predicted amino acid sequence of all 273 ORFs identified in bacteriophage Dp-1 that are identified as being expressed.

Table 31 shows the similarities identified in sequence between *Streptococcus* phage Dp-1 ORFs greater than 33 amino acids and sequences present in public sequence databases.

Table 32 shows the 4731 bp sequence of Dp-1 published by Sheehan et al., (1997).

Table 33 lists *Streptococcus pneumoniae* sequences listed in GenBank providing possible target sequences for inhibitory *Streptococcus pneumoniae* bacteriophage Dp-1 ORFs and other compounds with antibacterial activity

Background:

As indicated above, the present invention is concerned, in part, with the use of bacteriophage coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents. Thus, the invention concerns the selection of relevant bacteria. Particularly relevant bacteria are those which are pathogens of a complex organism such as an animal, e.g., mammals,

reptiles, and birds, and plants. Examples include *Staphylococcus aureus*, *Enterococcus* species, and *Streptococcus pneumoniae*. However, the invention can be applied to any bacterium (whether pathogenic or not) for which bacteriophage are available or which are found to have cellular components closely homologous to components targeted by phage of another bacterium.

Thus, the invention also concerns the bacteriophage which can infect a selected bacterium. Identification of ORFs or products from the phage which inhibit the host bacterium both provides an inhibitor compound and allows identification of the bacterial target affected by the phage-encoded inhibitor. Such targets are thus identified as potential targets for development of other antibacterial agents or inhibitors and the use of those targets to inhibit those bacteria. As indicated above, even if such a target is not initially identified in a particular bacterium, such a target can still be identified if a homologous target is identified in another bacterium. Usually, but not necessarily, such another bacterium would be a genetically closely related bacterium. Indeed, in some cases, a phage-encoded inhibitor can also inhibit such a homologous bacterial cellular component.

The demonstration that bacteriophage have adapted to inhibiting a host bacterium by acting on a particular cellular component or target provides a strong indication that that component is an appropriate target for developing and using antibacterial agents, *e.g.*, in therapeutic treatments. Thus, the present invention provides additional guidance over mere identification of bacterial essential genes, as the present invention also provides an indication of accessibility of the target to an inhibitor, and an indication that the target is sufficiently stable over time (*e.g.*, not subject to high rates of mutation) as phage acting on that target were able to develop and persist. Thus, the present invention identifies a subset of essential cellular components which are particularly likely to be appropriate targets for development of antibacterial agents.

The invention also, therefore, concerns the development or identification of inhibitors of bacteria, in addition to the phage-encoded inhibitory proteins (or RNA transcripts), which are active on the targets of bacteriophage-encoded inhibitors. As described herein, such inhibitors can be of a variety of different types, but are preferably small molecules.

The following description provides preferred methods for use in the various aspects of the invention. However, as those skilled in the art will readily recognize, other approaches can be used to obtain and process relevant information. Thus the invention is not limited to the specifically described methods. In addition, the following description provides a set of steps in a particular order. That series of steps

describes the overall development involved in the present invention. However, it is clear that individual steps or portions of steps may be usefully practiced separately, and, further, that certain steps may be performed in a different order or even bypassed if appropriate information is already available or is provided by other sources or methods.

Selecting and Growing Phage, and Isolating DNA

Conceptually, the first step involves selecting bacterial hosts of interest.

Preferably, but not necessarily, such hosts will be pathogens of clinical importance.

Alternatively, because bacteria all share certain fundamental metabolic and structural features, these features can be targeted for study in one strain, for example a nonpathogenic one, and extrapolated to similarly succeed in pathogenic ones. Nonpathogenic strains may also exhibit initial advantages in being not only less dangerous, but also, for example, in having better growth and culturing characteristics and/or better developed molecular biology techniques and reagents. Consequently, advantageously the invention provides the ability target virtually any bacteria, but preferably pathogenic bacteria, with antimicrobial compounds designed and/or developed using bacteriophage inhibitory proteins and peptides from phage with non-pathogenic and/or pathogenic hosts.

We have selected *Staphylococcus aureus*, *Streptococcus pneumoniae*, various *Enterococci*, and *Pseudomonas aeruginosa* as initial exemplary pathogens. These bacteria are a major cause of morbidity and mortality in hospital-based infections, and the appearance of antibiotics resistance in all three organisms makes it increasingly difficult to treat benign infections involving these organisms. Such infections can include, for example, otitis media, sinusitis, and skin, and airway infections (Neu, H.C. (1992). *Science* 257, 1064-1073). However, the approach described below is clearly applicable to any human bacterial pathogens including but not restricted to *Mycobacterium tuberculosis*, *Nisseria gonorrhoeae*, *Haemophilus influenza*, *Acinobacter*, *Escherichia coli*, *Shigella dysenteria*, *Streptococcus pyogenes*, *Helicobacter pylori*, and *Mycoplasma* species. This invention can also be applied to the discovery of anti-bacterial compounds directed against pathogens of animals other than humans, for example, sheep, cattle, swine, dogs, cats, birds, and reptiles. Similarly, the invention is not limited to animals, but also applies to plants and plant pathogens.

In general, the bacteria are grown according to standard methodologies employed in the art, including solid, semi-solid or liquid culturing, which procedures can be found in or extrapolated from standard sources such as Maloy, S.R., Stewart,

V.J., and Taylor, R.K. Genetic Analysis of Pathogenic Bacteria (1996) Cold Spring Harbor Laboratory Press, or Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; or Ausubel, F.M. et al. (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J. Culture conditions are selected which are adapted to the particular bacterium generally using culture conditions known in the art as appropriate, or adaptations of those conditions.

Nucleic acids within these bacteria can be routinely extracted through common procedures such as described in the above-referenced manuals and as generally known to those skilled in the art. Those nucleic acid stocks can then be used to practice the other inventive aspects described below.

Selection and Growth of Bacteriophage, and Isolation of DNA

The second step involves assembling a group of bacteriophages (phage collection) for one or more of the targeted bacterial hosts. While the invention can be utilized with a single bacteriophage for a pathogen or other bacterium, it is preferable to utilize a plurality of phage for each bacterium, as comparisons between a plurality of such phage provides useful additional information. Non-limiting examples of phage and sources for some of the above-mentioned pathogenic bacteria are found in Table 1. The criteria used to select such phages is that they are infectious for the microbe targeted, and replicate in, lyse, or otherwise inhibit growth of the bacterium in a measurable fashion. These phages can be very different from one another (representing different families), as judged by criteria such as morphology (head, tail, plate, etc.), and similarity of genome nucleotide sequence (cross-hybridization). Since such diverse bacteriophages are expected to block bacterial host metabolism and ultimately inhibit by a variety of mechanisms, their combined study will lead to the identification of different mechanisms by which the phages independently inhibit bacterial targets. Examples include degradation of host DNA (Parson K.A., and Snustad, D.P. (1975). *J. Virol.* 15, 221-444) and inhibition of host RNA transcription (Severinova, E., Severinov, K. and Darst, S.A. (1998). *J.Mol. Biol.* 279, 9-18). This, in turn, yields novel information on phage proteins that can inhibit the targeted microbe. As explained below, this 1) forms the basis of novel drug discovery efforts based on knowledge of the primary amino acid sequence of the phage inhibitor protein (e.g., peptide fragments or peptidomimetics) and/or 2) leads to the identification of bacterial biochemical pathways, the proteins of which are essential or significant for survival of the targeted microbe, and which enzymatic steps or

chemical reactions can be targeted by classical drug discovery methods using molecular inhibitors, for example, small molecule inhibitors.

Bacteriophage are generally either of two types, lytic or filamentous, meaning they either outright destroy their host and seek out new hosts after replication, or else continuously propagate and extrude progeny phage from the same host without destroying it. Regardless of the phage life cycle and type, preferred embodiments incorporate phage which impede cell growth in measurable fashion and preferably stop cell growth. To this end, lytic phage are preferred, although certain nonlytic species may also suffice, *e.g.*, if sufficiently bacteriostatic.

Various procedures that are commonly understood by those of skill in the art can be routinely employed to grow, isolate, and purify phage. Such procedures are exemplified by those found in such common laboratory aids such as Maloy, S.R., Stewart, V.J., and Taylor, R.K. Genetic Analysis of Pathogenic Bacteria (1996) Cold Spring Harbor Laboratory Press; Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; and Ausubel, F.M. et al. (eds.) (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J. The techniques generally involve the culturing of infected bacterial cells that are lysed naturally and/or chemically assisted, for example, by the use of an organic solvent such as chloroform that destroys the host cells thereby liberating the phage within. Following this, the cellular debris is centrifuged away from the supernatant containing the phage particles, and the phage then subsequently and selectively precipitated out of the supernatant using various methods usually employing the use of alcohols and/or other chemical compounds such as polyethylene glycol (PEG). The resulting phage can be further purified using various density gradient/centrifugation methodologies. The resulting phage are then chemically lysed, thereby releasing their nucleic acids that can be conveniently precipitated out of the supernatant to yield a viral nucleic acid supply of the phage of interest.

Exemplary bacteriophage are indicated in Table 1, along with sources where those phage may be obtained.

Exemplary bacteria include the reference bacteria for the identified bacteriophage, available from the same sources.

Characterizing Bacteriophage Genomes for ORFs

The third step involves systematically characterizing the genetic information contained in the phage genome. Within this genetic information is the sequence of all RNAs and proteins encoded by the phage, including those that are essential or

instrumental in inhibiting their host. This characterization is preferably done in a systematic fashion. For example, this can be done by first isolating high molecular weight genomic DNA from the phage using standard bacterial lysis methods, followed by phage purification using density gradient ultracentrifugation, and extraction of
5 nucleic acid from the purified phage preparation. The high molecular weight DNA is then analyzed to determine its size and to evaluate a proper strategy for its sequencing. The DNA is broken down into smaller size fragments by sonication or partial digestion with frequently cutting restriction enzymes such as Sau3A to yield predominantly 1 to 2 kilobase length DNA, which DNA can then be resolved by gel
10 electrophoresis followed by extraction from the gel.

The ends of the fragments are enzymatically treated to render them suitable for cloning and the pools of fragments are cloned in a bacterial plasmid to generate a library of the phage genome. Several hundred of these random DNA fragments contained in the plasmid vector are isolated as clones after introduction into an
15 appropriate bacterium, usually *Escherichia coli*. They are then individually expanded in culture and the DNA from each individual clone is purified. The nucleotide sequences of the inserts of these clones are determined by standard automated or manual methods, using oligonucleotide primers located on either side of the cloning site to direct polymerase mediated sequencing (e.g., the Sanger sequencing method or
20 a modification of that method). Other sequencing methods can also be used.

The sequence of individual clones is then deposited in a computer, and specific software programs (for example, Sequencher™, Gene Codes Corp.) are used to look for overlap between the various sequences, resulting in ordering of contig sequences and ultimately providing the complete sequence of the entire bacteriophage
25 genome (one such example is given in Table 2 for *Staphylococcus aureus* bacteriophage 77; others are also provided herein). This complete nucleotide sequence is preferably determined with a redundancy of at least 3- to 5-fold (number of independent sequencing events covering the same region) in order to minimize sequencing errors.

30 Preferably, the bacterial strain used as a phage host should not possess any other innate plasmids, transposons, or other phage or incompatible sequences that would complicate or otherwise make the various manipulations and analyses more difficult.

Commercially available computer software programs are used to translate the
35 nucleotide sequence of the phage to identify all protein sequences encoded by the phage (hereafter called open reading frames or ORFs). (Customized software can clearly also be used.) As phages are known to transcribe their genome into RNA from

both strands, in both directions, and sometimes in more than one frame for the same sequence, this exercise is done for both strands and in all six possible reading frames. As evolutionary constraints have forced the phage to conserve all of its vital protein sequences in as small a genome as possible, it is straightforward to identify all the proteins encoded by the phage by simple examination of the 6 translation frames of the genome. Once these ORFs are identified, they are cataloged into a phage proteome database (Table 3 lists ORFs identified from phage 77; ORF lists are also provided for other exemplary phage). This analysis is preferably performed for each phage under study. The process of ORF identification can be varied depending on the desired results. For example, the minimum length for the putative encoded polypeptide can be varied, and/or putative coding regions that have an associated Shine-Dalgarno sequence can be selected. In the case of phage 77 ORFs, such parameter adjustment was performed and resulted in the identification of ORFs as listed herein. Different parameters had resulted in the identification of the ORFs listed in the preceding U.S. Provisional Application 60/110,992, filed December 3, 1998, which is hereby incorporated by reference in its entirety.

Exemplary phage 77 ORFs identified in that provisional application and as identified herein are shown in the following table:

ORF ID from 60/110,992	Genomic position	a.a. size	Start codon	ORF ID from 241/190	Genomic position	a.a. size	Start codon
77ORF016	2369-24024	251	TTG	77ORF017	23269-23982	237	ATG
77ORF019	39845-40501	218	ATA	77ORF019	39851-40501	216	ATG
77ORF050	29268-29564	98	ATG	77ORF182	29268-29564	98	ATG
77ORF050	29268-29564	98	ATG	77ORF043	29304-29564	86	ATG
77ORF067	34312-34551	79	CTG	77ORF104	34393-34551	52	ATG
77ORF146	29051-29212	53	ATG	77ORF102	29051-29212	53	ATG

Identifying and Characterizing Inhibitory Phage ORFs

The fourth step entails identifying the phage protein or proteins or RNA transcripts that have the ability to inhibit their bacterial hosts. This can be accomplished, for example, by either or both of two non-mutually exclusive methods. The first method makes use of bioinformatics. Over the past few years, a large amount of nucleotide sequence information and corresponding translated products have become available through large genome sequencing projects for a variety of organisms including mammals, insects, plants, unicellular eukaryotes (yeast and fungi), as well as several bacterial genomes such as *E. coli*, *Mycobacterium tuberculosis*, *Bacillus subtilis*, *Staphylococcus aureus* and many others. Such sequences have been deposited in public databases (for example, non-redundant

sequence database at GenBank and SwissProt protein sequence database) (<http://www.ncbi.nlm.nih.gov>) and can be freely accessed to compare any specific query sequence to those present in such databases. For example, GenBank contains over 1.6 billion nucleotides corresponding to 2.3 million sequence records. Several
5 computer programs and servers (*e.g.*, TBLASTN) have been created to allow the rapid identification of homology between any given sequence from one organism to that of another present in such databases, and such programs are public and available free of charge.

In addition, it has been well established that basic biochemical pathways can
10 be conserved in very distant organisms (for example bacteria and man), and that the proteins performing the various enzymatic steps in these pathways are themselves conserved at the amino acid sequence level. Thus, proteins performing similar functions (*e.g.* DNA repair, RNA transcription, RNA translation) have frequently preserved key structural signatures, identifiable by similarities across regions of
15 proteins (domains and motifs). The antimicrobials of the present invention will preferably target features and targets that are highly characteristic or conserved in microbes, and not higher organisms.

Most genomes encode individual proteins or groups of proteins that can be assembled into protein families that have been evolutionarily conserved. Therefore,
20 similarity between a new query sequence and that of a member of a protein family (reference sequences from public databases) can immediately suggest a biochemical function for the novel query sequence, which in our case is a phage ORF.

The sequence homology between individual members of evolutionarily distant members of a protein family is usually not randomly distributed along the entire
25 length of the sequence but is often clustered into "motifs" and "domains". These correspond to key three-dimensional folds that form key catalytic and/or regulatory structures that perform key biochemical function(s) for the group of proteins. Commercially available computer software programs can identify such motifs in a new query sequence, again providing functional information for the query sequence.
30 Such structural and functional motifs have also been derived from the combined analysis of primary sequence databases (protein sequences) and protein structure databases (X-ray crystallography, nuclear magnetic resonance) using so-called "threading" methods (Rost B, and Sander C. (1996). *Ann. Rev. Biophys. Biomol. Struct.* 25, 113-136).

35 Such motifs and folds are themselves deposited in public databases which can be directly accessed (for example, SwissProt database; 3D-ALI at EMBL, Heidelberg; PROSITE). This basic exercise leads to a structural homology map in which each of

the phage ORFs has been probed for such similarities, and where initial structural and functional hits are identified (selected examples of sequence homologies detected between individual ORFs from the genome of *Staphylococcus aureus* bacteriophage 77 and sequences deposited in public databases are shown in Table 5 for ORFs 17/19/43/102/104/182).

This analysis can point out phage proteins with similarity to proteins from other phages (such as those for *E. coli*) playing an important role in the basic biochemical pathways of the phage (such as DNA replication, RNA transcription, tRNAs, coat protein and assembly). Selected examples of such proteins include integrase and capsid protein. Therefore, this analysis enables identification and elimination of non-essential ORFs as candidates for an inhibitor function, as well as the identification of (potentially) useful ones.

In addition, this analysis can point out specific ORFs as possible inhibitor ORFs. For example these ORFs may encode proteins or enzymes that alter bacterial cell structure, metabolism or physiology, and ultimately viability. Examples of such proteins present in the genome of *Staphylococcus aureus* bacteriophage 77 include orf14 (deoxyuridine triphosphatase from bacteriophage T5), and orf15 (sialidase). (These ORF identifications are as listed in provisional application 60/110,992.) Other examples include ORFs 9 and 12 of *S. aureus* phage 44 AHJD, which encode the putative lysis functions found in many bacteriophages – a “holin” and an “amidase”.

In addition, it is well known that bacterial and eukaryotic viruses can usurp pathways from their host in order to use them to their advantage in blocking host cellular pathways upon infection. The phage can achieve this by 1) directly producing an inhibitor of a key host pathway (e.g. T7 gene 0.5 and 2), 2) directly producing a novel activity (e.g. T4 DNA polymerase), and 3) altering concentrations of cell components by producing similar functions (e.g. T4 transfer RNAs). The identification of sequence similarity between phage ORFs and bacterial host genome sequences will be highly indicative of such a mechanism. (Selected examples of such homologies are listed in Figure 4 of the provisional application 60/110,992 and include orf4 (homologous to autolysin), orf20 (hypothetical protein from *Staphylococcus aureus*) and orf29 (hypothetical protein from *Staphylococcus aureus*)). These ORFs can be analyzed by a standard biochemical approach to directly test their inhibitor functions (e.g., as described below).

Alternatively, a homology search may reveal that a given phage ORF is related to a protein present in the databases having an activity known to be inhibitory, (e.g. inhibitor of host RNA polymerase by *E. coli* bacteriophage T7. Such a finding would implicate the phage ORF product in a related activity. This will also suggest that a

new antimicrobial could be derived by a mimetic approach (e.g., peptidomimetic) imitating this function or by a small molecule inhibitor to the bacterial target of the phage ORF, or any steps in the relevant host metabolic pathway, e.g., high throughput screening of small molecule libraries. Selected examples of such similarity between
5 ORFs of *Staphylococcus aureus* bacteriophage 77 and proteins with inhibitor functions for bacterial hosts are listed in Figure 4 of the provisional application 60/110,992. These include orf9 (similar to bacteriophage P1 *kilA* function), and orf4 (autolysin of *Staphylococcus aureus*, amidase enzymatic activity).

A reason for the biochemical study of individual ORFs for inhibitor function is
10 that their expression or overexpression will block cellular pathways of the host, ultimately leading to arrest and/or inhibition of host metabolism. In addition, such ORFs can alter host metabolism in different ways, including modification of pathogenicity. Therefore, individual ORFs identified above are expressed, preferably overexpressed, in the host and the effect of this expression or overexpression on host
15 metabolism and viability is measured. This approach can be systematically applied to every ORF of the phage, if necessary, and does not rely on the absolute identification of candidate ORFs by bioinformatics. Individual ORFs are resynthesized from the phage genomic DNA, e.g., by the polymerase chain reaction (PCR), preferably using oligonucleotide primers flanking the ORF on either side. These single ORFs are
20 preferably engineered so that they contain appropriate cloning sites at their extremities to allow their introduction into a new bacterial expression plasmid, allowing propagation in a standard bacterial host such as *E. coli*, but containing the necessary information for plasmid replication in the target microbe such as *S. aureus* (hereafter referred to as shuttle vector). Shuttle vectors and their use are well known in the art.

25 Such shuttle vectors preferably also contain regulatory sequences that allow inducible expression of the introduced ORF. As the candidate ORF may encode an inhibitor function that will eliminate the host, it is beneficial that it not be expressed prior to testing for activity. Thus, screening for such sequences when expressed in a constitutive fashion is less likely to be successful when the inhibitor is lethal. In the
30 exemplary inducible system presented in Figure 1A, 1B, 2, and 7, regulatory sequences from the *ars* operon of *S. aureus* are used to direct individual ORF expression in *S. aureus* (or other bacteria in which the *ars* system is functional). The *ars* operon encodes a series of proteins which normally mediate the extrusion of arsenite and other trivalent oxyanions from the cells when they are exposed to such
35 toxic substances in their environment. The operon encoding this detoxifying mechanism is normally silent and only induced when arsenite-related compounds are

present. (Tauriainen, S. et al. (1997) *App. Env. Microb.*, Vol. 63, No. 11, p. 4456-4461.)

Therefore, individual phage ORFs can be expressed in *S. aureus* in an inducible fashion by adding to the culture medium non-toxic arsenite concentrations during the growth of individual *S. aureus* clones expressing such individual phage ORFs. Toxicity of the phage inhibitor ORF for the host is monitored by reduction or arrest of growth under induction conditions, as measured by optical density in liquid culture or after plating the induced cultures on solid medium. Subsequently, interference of the phage ORF with the host biochemical pathways ultimately leading to reduced or arrested host metabolism can be measured by pulse-chase experiments using radiolabeled precursors of either DNA replication, RNA transcription, or protein synthesis. Similar constructs can be made and used for other bacteria using well-known techniques.

Those skilled in the art are familiar with a variety of other inducible systems which can also be used for the controlled expression of phage ORFs, including, for example, lactose (see *e.g.*, Stratagene's LacSwitchTMII system; La Jolla, CA) and tetracycline-based systems (see, *e.g.* Clontech's Tet On/Tet OffTM system; Palo Alto, CA). The arsenite-inducible system described is further depicted in Figures 1, 2 and 7.

The selection or construction of shuttle vectors and the selection and use of inducible systems are well known and thus other shuttle vectors appropriate for other bacteria can be readily provided by those skilled in the art, *e.g.*, for use in other bacterial species.

Standard methodologies for expressing proteins from constructs, and isolating and manipulating those proteins, for example in cross-linking and affinity chromatography studies, may be found in various commonly available and known laboratory manuals. See, *e.g.*, Current Protocols in Protein Science, John Wiley & Sons, Secaucus, N.J., and Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.

It has been found that certain phage or other viruses inhibit host cells, at least in part, by producing an antisense RNA which binds to and inhibits translation from a bacterial RNA sequence. Thus, in the case of potentially inhibitor RNA transcripts encoded by the phage genome, a strong indicator of a possible inhibitory function is provided by the identification of phage sequence which is the identical to or fully complementary (or with only a small percentage of mismatch, *e.g.*, <10%, preferably less than 5%, most preferably less than 3%, to a bacterial sequence. This approach is convenient in the case of bacteria that have been essentially completely sequenced, as the comparison can be performed by computer using public database information.

The inhibitory effect of the transcript can be confirmed using expression of the phage sequence in a host bacterium. If needed, such inhibitory can also be tested by transfecting the cells with a vector that will transcribe the phage sequence to form RNA in such manner that the RNA produced will not be translated into a polypeptide.

5 Inhibition under such conditions provides a strong indication that the inhibition is due to the transcript rather than to an encoded polypeptide.

In an alternative, the expression of an ORF in a host bacterium is found to be inhibitory, but the inhibition is found to be due to an RNA product of the genomic coding region. For antisense inhibition, the sequence of the bacterial target nucleic acid sequence can be identified by inspection of the phage sequence, and the full

10 sequence of the relevant coding region for the bacterial product can be found from a database of the bacterial genomic sequence or can be isolated by standard techniques (e.g., a clone in a genomic library can be isolated which contains the full bacterial ORF, and then sequenced).

15 In either case, the identification of a target which is inhibited by an RNA transcript produced by a phage provides both the possible inhibition of bacteria naturally containing the same target nucleic acid sequence, as well as the ability to use the target sequence in screening for other types of compounds which will act directly on the target nucleic acid sequence or on a polypeptide product expressed or

20 regulated, at least in part, by the target of the inhibitory phage RNA.

In some cases it will be found that the target of an inhibitory phage RNA or protein has previously been found to be a target of an inhibitory phage RNA or protein has previously been found to be a target for an antibacterial agent. In such cases, the phage inhibitor can still provide useful information if it is found that the

25 phage-encoded product acts at a different site than the previously identified antibacterial agent or inhibitor, *i.e.*, acts at a phage-specific site. For many targets, action at a different site provides highly beneficial characteristics and/or information. For example, an alternate site of inhibitor action can at least partially overcome a resistance mechanism in a bacterium. As an illustration, in many cases, resistance is

30 due, in large part, to altered binding characteristics of the immediate target to the antibacterial agent. The altered binding is due to a structural change which prevents or destabilizes the binding. However, the structural change is frequently quite local, so that compounds which bind at different local sites will be unaffected or affected to a much lesser degree. Indeed, in some cases the local sites will be on a different

35 molecule and so may be completely unaffected by the local structural change creating resistance to the original agent(s). An example of resistance due to altered binding is

provided by methicillin-resistant *Staphylococcus aureus*, in which the resistance is due to an altered penicillin-binding protein.

In other cases, a new site of action can have improved accessibility as compared to a site acted on by a previously identified agent. This can, for example, assist in allowing effective treatment at lower doses, or in allowing access by a larger range of types of compounds, potentially allowing identification of more potential active agents.

Another advantage is that the structural characteristics of a different site of action will lead to identification and/or development of inhibitors with different structures and different pharmacological parameter. This can allow a greater range of possibilities when selecting an antibacterial agent.

Yet further, different sites often produce different inhibitory characteristics in the target organism. This is commonly the case for multi-domain target proteins. Thus, inhibition targeting an alternate site can produce more efficacious action, e.g., faster killing, slower development of resistance, lower numbers of surviving cells, and different secondary effects (for example, different nutrient utilization).

Staphylococcus aureus phage 77

As indicated above, the present invention is concerned, in part, with the use of bacteriophage 77 coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents.

As described, phage 77 ORFs 17, 19, 43, 102, 104, and 182 have been found to have bacteria inhibiting function. Identification of ORFs 17, 19, 43, 102, 104, and 182 and products from the phage which inhibit the host bacterium both provides an inhibitor compound and allows identification of the bacterial target affected by the phage-encoded inhibitor. Such a target is thus identified as a potential target for development of other antibacterial agents or inhibitors and the use of those targets to inhibit those bacteria. As indicated above, even if such a target is not initially identified in a particular bacterium, such a target can still be identified if a homologous target is identified in another bacterium. Usually, but not necessarily, such another bacterium would be a genetically closely related bacterium. Indeed, in some cases, an inhibitor encoded by phage 77 ORF 17, 19, 43, 102, 104, or 182 can also inhibit such a homologous bacterial cellular component.

Possible bacterial target sequences are described herein by reference to sequence source sites. In preferred embodiments, the sequence encoding the target corresponds

to a *S. aureus* nucleic acid sequence available from numerous sources including *S. aureus* sequences deposited in GenBank, *S. aureus* sequences found in European Patent Application No. 97100110.7 to Human Genome Sciences, Inc. filed January 7, 1997, *S. aureus* sequences available from TIGR at

- 5 <http://www.tigr.org/tdb/mdb/mdb.html>, and *S. aureus* sequences available from the Oklahoma University *S. aureus* sequencing project at the following URL: http://www.genome.ou.edu/staph_new.html. Such possible targets are particularly applicable to *S. aureus* phages 77, 3A, 96, and 44 AHJD.

- 10 The amino acid sequence of a polypeptide target is readily provided by translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. Also, in preferred embodiments, a target sequence corresponds to a *S. aureus* coding sequence corresponding to a sequence listed in Table 15 herein. The listing in Table 15 describes *S. aureus* sequences currently listed with GenBank. Again, for the sake of brevity, the sequences are described by
- 15 reference to the database accession numbers instead of being written out in full herein. In cases where an entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a phage host *S. aureus* genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional
- 20 sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

Staphylococcus aureus phage 44 AHJD

- 25 The present invention also can utilize the identification of naturally occurring DNA sequence elements within *Staphylococcus aureus* bacteriophage 44AHJD which encode proteins with antimicrobial activity.

- Such identification can utilize bioinformatics identification of specific proteins (ORFs) utilized by *Staphylococcus aureus* bacteriophage 44AHJD during the viral life
- 30 cycle, resulting in a slowing or arrest of growth of the bacterial host, or in death, of the *Staphylococcus aureus* host including lysis of the infected bacteria. Thus, some of the bacteriophage 44AHJD DNA sequences encoding these proteins (ORFs) are predicted to encode antimicrobial functions. Information derived from these DNA sequences and translated ORFs can, in turn, be utilized to develop inhibitory
- 35 compounds by peptidomimetics that can also function as antimicrobials. In addition, the identification of the host bacterial proteins that are targeted and inhibited by the

antimicrobial bacteriophage ORFs can themselves provide novel targets for drug discovery.

The methodology described above is used to identify and characterize DNA sequences from *Staphylococcus* sp. bacteriophage 44 AHJD that have antimicrobial activity. As described in the Examples, the *Staphylococcus aureus* propagating strain (PS 44A), obtained from the Felix d'Herelle Reference Centre (#HER 1101), was used as a host to propagate its phage 44AHJD, also obtained from the Felix d'Herelle Reference Centre (#HER 101). By sequencing, we found that bacteriophage 44AHJD consists of 16,668 bp (Table 16) predicted to encode 73 ORFs greater than 33 amino acids (Tables 17 & 18). Computational analysis of the predicted protein products of *Staphylococcus aureus* bacteriophage 44AHJD identified homolgs in public sequence databases as listed in Table 19 and 20, along with the accompanying list of related proteins.

From this analysis, it is apparent that 3 genes (ORF 3, 7, and 8) are related to structural proteins found in other bacteriophages. These include genes predicted to encode a tail protein (ORF 3), an upper collar/connector protein of the phage virion (ORF 7), and a lower collar protein (ORF 8). Bioinformatics has also identified one gene whose product is likely involved in phage DNA synthesis. One gene (ORF 1) shows significant homology to DNA polymerases of a number of bacteriophages, bacteria and fungi, and the product of this gene is likely responsible for replicating the genetic material of bacteriophage 44AHJD. ORF 2 encodes a protein with homology to the *dinC* gene of *Bacillus subtilis* that encodes a protein involved in teichoic acid biosynthesis. Teichoic acid is a polyphosphate polymer found in some, but not all, Gram positive organisms (and not in Gram negative organisms), where it is attached to the peptidoglycan layer. The phage protein may thus be involved in the synthesis of this material for incorporation into the cell wall, allowing enhanced lysis by the phage lysis enzymes or, as many enzymes can function in "reverse reactions", may be involved in its degradation allowing for penetration of the peptidoglycan and phage genome entry into the cell following adsorption. The similarity between *Staphylococcus aureus* bacteriophage 44AHJD and *E. coli* phage T7 indicates that they may share similar mechanisms of replication and growth. Both phages belong to the Pododviridae Family of bacteriophages and are members of the "T7-like" Genus of this Family (Ackermann and DuBow; VIth ICTV Report).

Two genes, ORF 9 and 12, were identified with the potential to encode antimicrobial protein products. The homology alignments are shown in Tables 19 and 20. The predicted product of ORF 9 is related to a class of genes which encodes lysozyme-like functions, enzymes which cleave linkages in the mucopolysaccharide cell wall structure of a variety of micro-organisms, including that from the *Staphylococcus aureus* bacteriophage Twort. ORF 12 of *Staphylococcus aureus* bacteriophage 44AHJD shows homology to a set of lysis proteins from several bacteriophages. These lysis proteins are also referred to as holins, and represent phage-encoded lysis functions required for transit of the phage murein hydrolases (lysozyme) to the periplasm, where it can digest the cell wall and thus lyse the bacterium.

Thus, in particular embodiments, the present invention provides a nucleic acid sequence isolated from *Staphylococcus aureus* bacteriophage 44AHJD comprising at least a portion of one of the genes described above with antimicrobial activity. For example, ORF 1 encodes a DNA polymerase function. This polymerase may utilize host-derived accessory proteins for its activity when replicating the phage template, sequestering such proteins from use by the bacterial polymerase, resulting in inhibition of DNA replication, cell division, and cell growth. Alternatively, ORF 9 directly encodes a polypeptide with antimicrobial activity. ORF 9 is predicted to encode an amidase, a protein known to act as a cell wall degrading enzyme. ORF 12 likely encodes a holin function required for transit of the phage amidase (gene 9 product) to the periplasm. When this type of gene product from Bacillus phage phi 29 (gene 14), was cloned in *Escherichia coli*, cell death ensued (Steiner et al., 1993). Thus, production of proteins from Bacillus phage phi 29 gene 14 in *E. coli* resulted in cell death, whereas production of protein from Bacillus phage phi 29 gene 14 concomitantly with the phi 29 lysozyme or unrelated murein-degrading enzymes led to lysis, suggesting that membrane-bound protein 14 induces a nonspecific lesion in the cytoplasmic membrane (Steiner et al., 1993).

The present invention also provides the use of the *Staphylococcus* bacteriophage 44 AHJD antimicrobial ORFs or ORF products as pharmacological agents, either wholly or in part and derivatives, as well as the use of corresponding peptidomimetics, developed from amino acid or nucleotide sequence knowledge derived from *Staphylococcus* bacteriophage 44 AHJD killer ORFs.

Enterococcus phage 182

Bacteriophage 182 was obtained from the Felix D'Herelle phage collection (Ste. Foy, Quebec) and infects *Enterococcus* sp. Group D. The genome of
5 *Enterococcus* bacteriophage 182 consists of 17,833 bp (Table 21) and is predicted to encode 80 ORFs greater than 33 amino acids (Tables 22 and 23). Computational analysis of the predicted protein products of *Enterococcus* bacteriophage 182 was performed in order to identify protein products related to those deposited in public databases. Bacteriophage 182 protein products which detected sequences with
10 significant sequence similarity in public databases are listed in Table 24 and 26, along with the accompanying list of related proteins.

From this analysis, it is apparent that 5 genes (ORF 001, 004, 007, 009, and 011) are related to structural proteins of several *Bacillus* phages – *Bacillus* bacteriophage PZA, phi-29, and B103. These include genes predicted to encode a tail
15 protein (ORF 001), a head protein (ORF 004), and upper collar protein (ORF 007), a lower collar protein (ORF 009), and a pre-neck appendage protein (ORF 011). Two gene products are predicted to encode genes which direct phage morphogenesis – these are ORF 005 and 019.

Bioinformatics has also identified three genes whose products are likely
20 involved in phage DNA synthesis. One gene, ORF 002 shows significant homology to DNA polymerases of a number of bacteriophages, and the product of this gene is likely responsible for replicating the genetic material of bacteriophage 182. ORF 006 encodes a protein with homology to the encapsidation proteins of several other bacteriophages, including *Bacillus* phage phi-29 (P11014), PZA (P07541), and B103
25 (X99260) and *Streptococcus* phage CP-1 (Z47794). These gene products catalyze the *in vivo* and *in vitro* genome-encapsidation reaction (Garvey et al., 1985). Proteins involved in genome packaging have been shown to have additional activities that affect biochemical reactions in other phages and their hosts. For example, the coat protein of the RNA bacteriophage MS2 interacts with viral RNA to translationally
30 repress replicase synthesis (Pickett and Peabody, 1993). This protein-RNA interaction also plays a role in genome encapsidation, enveloping a single copy of the viral genome in a protein shell composed of many molecules of coat protein. In addition, the bacteriophage λ terminase enzyme can be lethal to *E. coli* when expressed,

suggesting cleavage of packaging sites in the bacterial chromosome. Also present within bacteriophage 182 is a gene, ORF 010, that encodes a protein that is related to the terminal proteins of *Bacillus* phage Nf (P06812), *Bacillus* phage GA-1 (X96987) and *Bacillus* phage B103 (X99260). DNA terminal proteins are linked to the 5' ends of both strands of the genome and are essential for DNA replication playing a role in initial priming of DNA replication. The similarity between *Enterococcus* bacteriophage 182 and *Bacillus* phages phi-29, PZA, and B103 indicates that they may share similar mechanisms of replication and growth. Protein-primed DNA replication is a well described phenomenon, and in the phi-29-like phages, the ends of the DNA serve as origins and termini of replication (Gutiérrez et al., 1986; Yoshikawa et al., 1985).

There is also a gene (ORF 015) that encodes a protein showing homology to an early protein product of *Bacillus* bacteriophage PZA and the single-strand nucleic acid binding protein of bacteriophage B103.

Two genes, ORF 008 and 014, were identified with the potential to encode anti-microbial protein products. The homology alignments are shown in Tables 24 & 26 and biochemical features of the predicted polypeptides shown in Table 25. The predicted product of ORF 008 is related to a class of genes which encodes lysozyme-like functions, enzymes which cleave linkages in the mucopolysaccharide cell wall structure of a variety of micro-organisms. ORF 014 of *Enterococcus* 182 shows homology to a set of lysis proteins from *Bacillus* bacteriophage phi-29, PZA, and B103. These lysis proteins are also referred to as holins and represent phage encoded lysis functions required for transit of the phage murein hydrolases (lysozyme) to the periplasm, where it can digest the outer cell wall and thus lyse the bacterium.

Thus, the present invention provides a nucleic acid sequence obtained from *Enterococcus* bacteriophage 182 comprising at least a portion of a phage 182 ORF, preferably an inhibitory ORF, and more preferably at least a portion of one of the genes described above with anti-microbial activity. For example, ORF 002 encodes a DNA polymerase function. This polymerase may utilize host-derived accessory proteins for its activity when replicating the phage template, sequestering such proteins from use by the bacterial polymerase, resulting in inhibition of DNA replication, cell division, and cell growth. Alternatively, ORFs 008 or 014 directly encode polypeptides with anti-microbial activity. ORF 008 is predicted to encode an

autolytic lysozyme, a protein known to have anti-microbial activity (Martin *et al.*, 1998). ORF 014 likely encodes a holin function required for transit of the phage murein hydrolases to the periplasm. When the related product from *Bacillus* phage phi 29 (gene 14), was cloned in *Escherichia coli*, cell death ensued (Steiner *et al.*, 1993).

5 Thus, production of proteins from *Bacillus* phage phi 29 gene 14 in *E. coli* resulted in cell death, whereas production of protein from *Bacillus* phage phi 29 gene 14 concomitantly with the phi 29 lysozyme or unrelated murein-degrading enzymes led to lysis, suggesting that membrane-bound protein 14 induces a nonspecific lesion in the cytoplasmic membrane (Steiner *et al.*, 1993).

10 The present invention also provides the use of the *Enterococcus* bacteriophage 182 anti-microbial ORFs as pharmacological agents, either wholly or in part and derivatives, as well as the use of corresponding peptidomimetics, developed from amino acid or nucleotide sequence knowledge derived from *Enterococcus* bacteriophage 182 killer ORFs. This can be done where the structure of the
15 peptidomimetic compound corresponds to the structure of the active portion of a product of an ORF. In this analysis, the peptide backbone is transformed into a carbon based hydrophobic structure that can retain cytostatic or cytotoxic activity for the bacterium. This is done by standard medicinal chemistry methods, measuring growth inhibition of the various molecules in liquid cultures or on solid medium. These
20 mimetics also represent lead compounds for the development of novel antibiotics. In this context, "corresponds" means that the peptidomimetic compound structure has sufficient similarities to the structure of the active portion of a product of one of the *Enterococcus* ORFs listed, that the peptidomimetic will interact with the same molecule as the product of the ORF, and preferably will elicit at least one cellular
25 response in common which relates to the inhibition of the cell by the phage protein.

To validate the identity of an ORF as a killer ORF, it is preferably expressed in the host or other test bacterial organism and the effect of this expression on bacterial growth and replication is assessed. Therefore, all individual ORFs identified herein, e.g., those identified above, can be expressed, preferably overexpressed, in a
30 suitable host bacterium e.g., a host *Enterococcus* and the effect of this expression or overexpression on host metabolism and viability can be measured.

Individual ORFs can be resynthesized from the phage genomic DNA by the polymerase chain reaction (PCR) using oligonucleotide primers flanking the ORF on

either side. Those skilled in the art are familiar with the design and synthesis of appropriate primer sequences. These single ORFs are preferably engineered so that they contain appropriate cloning sites at their extremities to allow their introduction into a new bacterial expression plasmid, allowing propagation in a standard bacterial host such as *E. coli*, but containing the necessary information for plasmid replication in the target microbe, *Enterococcus* sp. (hereafter referred to as a shuttle vector).

This shuttle vector also preferably contains regulatory sequences that allow inducible expression of the introduced ORF. As the candidate ORF may encode a killer function that will eliminate the host, it is highly advantageous that it not be expressed (or at least not expressed at a substantial level) prior to testing for activity; thus screening for such sequences in a constitutive fashion is less likely to be successful (lethality). In an example presented in Fig. 7, regulatory sequences from the *ars* operon are used to direct individual ORF expression in *Enterococcus*. The *ars* operon encodes a series of proteins which normally mediate the extrusion of arsenite and several other trivalent oxyanions from the cells when they are exposed to such toxic substances in their environment. The operon encoding this detoxifying mechanism is normally silent and only induced when arsenite-related compounds are present.

Therefore, individual phage ORFs can be expressed in *Enterococcus* or other suitable host in an inducible fashion by adding to the culture medium non-toxic arsenite concentrations during the growth of individual *Enterococcus* (or other host cells) clones expressing such individual phage ORFs. Toxicity of the phage killer ORF for the host is monitored by reduction or arrest of growth under induction conditions, as measured by optical density in liquid culture or after plating the induced cultures on solid medium. Subsequently, interference of the phage ORF with the host biochemical pathways ultimately leading to reducing or arresting host metabolism can be measured by pulse chase experiments using radiolabeled precursors of either DNA replication, RNA transcription, or protein synthesis.

Of course, other inducible regulatory sequences (e.g., promoters, operators, etc.) may be used (e.g., systems using positive induction of expression or systems using release of repression). A variety of such systems are known to those skilled in the art and can be utilized in the present invention.

Nucleic acid sequences of the present invention can be isolated using a method similar to those described herein or other methods known to those skilled in the art. In addition, such nucleic acid sequences can be chemically synthesized by well-known methods. Having the phage 182 ORFs, e.g., anti-bacterial ORFs of the present invention, portions thereof, or oligonucleotides derived therefrom as described, other anti-microbial sequences from other bacteriophage sources can be identified and isolated using methods described here or other methods, including methods utilizing nucleic acid hybridization and/or computer-based sequence alignment methods.

The invention also provides bacteriophage anti-microbial DNA segments from other phages based on nucleic acids and sequences hybridizing to the presently identified inhibitory ORF under high stringency conditions or sequences which are highly homologous. The bacteriophage anti-microbial DNA segment from bacteriophage 182 can be used to identify a related segment from another unrelated phage based on stringent conditions of hybridization or on being a homolog based on nucleic acid and/or amino acid sequence comparisons. As with the phage 182 inhibitory sequences, such homologous coding sequences and products can be used as antimicrobials, to construct active portions or derivatives, to construct peptidomimetics, and to identify bacterial targets.

Enterococcus sequences are listed in Table 27 by accession number, providing identification of possible targets of *Enterococcus* phage inhibitory ORF products, e.g., from phage 182.

Streptococcus pneumoniae

As indicated in the Summary above, the present invention is concerned with the use of *Streptococcus* sp. bacteriophage Dp-1 coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents.

Streptococcus pneumoniae is an important cause of community-acquired pneumonia and a major cause of otitis media, sinusitis, and meningitis in children and adults. In Spain and other Mediterranean countries, the majority of *S. pneumoniae* are relatively resistant to penicillin (Klugman, 1990; Fenoll et al., 1991; Jørgensen et al., 1990). These strains also have decreased susceptibility to broad-spectrum cephalosporins, which are frequently used in the empiric treatment of meningitis and

other serious invasive bacterial infections. High-level resistance of pneumococci has been encountered in Hungary where 70% of children who were colonized with *S. pneumoniae* carried penicillin resistant strains that were also resistant to tetracycline, erythromycin, trimethoprim/sulfamethoxazole, and 30% resistant to chloramphenicol (Neu, 1992). The resistance of pneumococci to macrolides such as erythromycin averages 20-25% in France, ~20% in Japan, and <10% in Spain (Neu, 1992).

The antimicrobial susceptibilities and distribution of serotypes of the 42 isolates of *S. pneumoniae* in southern Taiwan from invasive infections have been recently determined (Hseuh et al., 1996). Resistance rates among these isolates were: erythromycin, 61.9%; clindamycin, 47.6%; chloramphenicol, 19%; and tetracycline, 73.8%. Resistance to three or more classes of antibiotics was found in 33.3% of the isolates. Bacteremic pneumonia and primary bacteremia accounted for 64.3% of the infections and mortality was 42.6%. Given the severity of these infections despite adequate antibiotic therapy, there is clearly a need for introduction of new therapeutic options to prevent mortality due to invasive *S. pneumoniae* infections.

Pneumococcal phages belong to four families and they present a great variety in morphology, including lytic and temperate phages (for a review, see Garcia et al., 1997). Examples of lytic phages are Cp-1 and Dp-1, whereas examples of temperate phages are HB-3, EJ-1, and HB-746. The complete nucleotide sequence and functional organization of Cp-1 has been reported (Martin et al., 1996). Cp-1 has a 19,345 bp double-stranded DNA genome, with a terminal protein covalently linked to its 5' ends, that replicates by a protein primed mechanism. The phage contains 29 ORFs, 23 on one strand and 6 on the opposite. When these predicted proteins were compared to sequences compiled in GenBank EMBL databases, to ORFs showed significant similarity to proteins of bacteriophage 29 that infects *B. subtilis* (Martin et al., 1996). The similar proteins corresponded to those involved in DNA replication (terminal protein and DNA polymerase), structural and morphogenic proteins (major head, collar, connector, tail, and encapsidation proteins), and proteins involved in lysis function (holin and lysozyme). In its strategy of lysis, the holin gene product inserts itself into the cell membrane, allowing access of the lysozyme to the peptidoglycan. Expression of the Cp-1 holin protein in *E. coli* results in cell death after 2-hours of induction, but did not lead to lysis (Garcia et al., 1997). Cells harboring a plasmid construction with holin and lysozyme genes together did lyse after induction and the

viability loss was similar to that of the culture expressing holin alone. Cloning of these lytic genes in *S. pneumoniae* showed that both genes had the same effect as in *E. coli*. That is, holin itself did not lyse the culture but the viability loss was noticeable, whereas both holin and lysozyme together were capable of lysing M31, an amidase deleted mutant (Garcia et al., 1997).

Recently, a small portion (~4 kbp) of a second *S. pneumoniae* phage, Dp-1, has been sequenced (Sheehan et al., 1997). This portion contains the genes coding for the lytic system (Sheehan et al., 1997) and shows a modular organization similar to that described for Cp-1. However, in this case, a single chimeric protein appears to be made in which the N-terminal domain is highly similar to that of the murein hydrolase coded by a gene found in the phage BK5-T that infects *Lactococcus lactis*, and the C-terminal domain is homologous to holins. Thus, both functions appear to have been combined in a novel chimeric protein.

Bacteriophage Dp-1 was obtained from Dr. P. Garcia (Departamento de Microbiologia Molecular, Centro de Departamento de Investigaciones Biologicas, Consejo Superior de Investigaciones Cientificas, Velazquez, Madrid, Spain). We found that Dp-1 has a double-stranded DNA genome of 56,506 bp, predicted to encode 85 ORFs greater than 33 amino acids and with upstream Shine-Dalgarno motifs for translation initiation (Tables 28 & 30, and Fig. 6). Computational analysis of the predicted protein products of *Streptococcus* bacteriophage Dp-1 protein products, which detected homologs in public databases, are listed in Table 31, along with the accompanying list of related proteins.

From this analysis, it is apparent that several predicted genes of Dp-1 encode polypeptides that are related to structural proteins. ORFs 001, 002, 004, and 030 are predicted to encode tail proteins, minor structural proteins, and minor capsid proteins (Table 31). We also note the identification of several gene products that are likely involved in DNA synthesis. These include ORF 3 which encodes DNA polymerase, ORF 8 which encodes a SWI/SNF helicase-related protein, ORF 10 encodes a protein showing homology to recA, and ORF 13 encodes a dnaZX-like ORF.

In *E. coli*, RapA encodes an RNA polymerase (RNAP)-associated protein with ATPase activity and which is a homolog of the eukaryotic SWI/SNF family, a set of proteins whose members are involved in transcription activation, nucleosome remodeling, and DNA repair. RapA forms a stable complex with RNAP,

as if it were a subunit of RNAP and it is possible that the ORF 8 product behaves similarly or in a dominant-negative fashion to inhibit the activity of RapA. Mutation of the essential *E. coli* dnaZX results in a block in DNA chain elongation during replication (Maki et al., 1988). The dnaZX gene has only one open reading frame for
 5 a 71-kDa polypeptide from which the two distinct DNA polymerase III holoenzyme subunits, tau (71 kDa) and gamma (47 kDa), are produced. The tau subunit is the precursor of the gamma subunit, and the gamma subunit is produced by a -1 frameshift causing early termination of translation (Tsuchihashi et al., 1990). These proteins show single-strand DNA binding properties that is ATPase (and dATPase)
 10 dependent and are thought to increasing the processivity of the core DNA polymerase enzyme (Lee et al., 1987).

There are several Dp-1 ORFs which encode proteins predicted to play a role in cellular metabolic pathways. These include polypeptides involved in coenzyme PQQ synthesis (ORFs 20, 29, 38). Pyrrolo-quinoline quinone (PQQ) is the non-covalently
 15 bound prosthetic group of many quinoproteins catalysing reactions in the periplasm of Gram-negative bacteria. Most of these involve the oxidation of alcohols or aldose sugars. Interestingly, ORFs 20, 29, and 30 also show homology to the exoenzyme S regulon (Frank, 1997). Proteins encoded by the *P. aeruginosa* exoenzyme S regulon may be involved in a contact-mediated translocation mechanism to transfer anti-host
 20 factors directly into eukaryotic cells disrupting eukaryotic signal transduction through ADP-ribosylation (Frank, 1997).

There is also a protein with similarity to GTP cyclohydrolase I (ORF 21) and ORF 41 which shows homology to dUTPase (Table 31). GTP cyclohydrolase I is an enzyme that catalyzes the first reaction in the pathway for the biosynthesis of the
 25 pteridine, a cofactor of the monooxygenases of the aromatic amino acids. Disruption of the homologous gene in *Saccharomyces cerevisiae* leads to a recessive conditional lethality due to folinic acid auxotrophy, that can be complemented with the mammalian or bacterial GTP cyclohydrolase I enzymes (Nardese et al., 1996; Mancini et al., 1999).

30 ORF 16 shows high homology to autolysin. This region of the phage sequence was previously reported (Sheehan et al., 1997) and encompasses ~ 4 kbp of our sequence. The sequence published by (Sheehan et al., 1997) is shown in Table 32.

Thus, the present invention provides a nucleic acid sequence obtained from *Streptococcus* bacteriophage Dp-1 comprising at least a portion of a phage Dp-1 ORF;
 35 preferably an inhibitory ORF, and more preferably at least a portion of one of the genes described above with anti-microbial activity. For example, ORF 013 encodes a

protein with homology to the gamma subunit of DNA polymerase (dnaX gene). This protein may act in a dominant-negative fashion to sequester the host DNA polymerase for its own replication, thus inhibiting host DNA replication. The dnaX gene product is essential for *E. coli* replication (Kodaira et al., 1983).

5 In certain preferred embodiments of the present invention, the bacterial target of a bacteriophage inhibitor ORF product, e.g., an inhibitory protein or polypeptide, is encoded by a *Streptococcus* nucleic acid coding sequence from a host bacterium for bacteriophage Dp-1. As above, possible target sequences are described herein by reference to sequence source sites. The sequence encoding the target preferably
10 corresponds to a *Streptococcus* nucleic acid sequence available from The Institute for Genomic Research (TIGR), or available from GenBank or other public database. The TIGR *Streptococcus* sequences are publicly available at The Institute for Genomics Research at URL: <http://www.tigr.org>

The amino acid sequence of a polypeptide target is readily provided by
15 translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. Also, in preferred embodiments, a target sequence corresponds to a *Streptococcus pneumoniae* coding sequences corresponding to a sequence listed in Table 33 herein. Sequences for other Streptococcal species are also available from TIGR and/or from GenBank. The listing in Table 33 describes
20 *Streptococcus* sequences currently deposited in GenBank. Again, for the sake of brevity, the sequences are described by reference to the GenBank entries instead of being written out in full herein. In cases where the TIGR or GenBank entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a phage Dp-1 host *Streptococcus* sp.
25 genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

30 In the various aspects of this invention involving Dp-1 sequences, preferably the sequence is preferably not contained in the sequence described in Sheehan et al., 1997 (Table 32).

Validating Identified Inhibitory Phage ORFs

35 A fifth step involves validating the identified phage inhibitor ORF by independent methods, and delineating further possible smaller segments of the ORFs

that have inhibitory activity. Several methods exist to validate the role of the identified ORF as an inhibitor ORF.

One example utilizes the creation of a mutant variant of the phage ORF in which the candidate ORF carries a partial or complete loss-of-function mutation that is measurable as compared with the non-mutant ORF. Comparison of the effects of expression of the loss of function mutant with the normal ORF provides confirmation of the identification of an inhibitor ORF where the loss-of-function mutant provides a measurably lower level of inhibition, preferably no inhibition. The loss of function may be conditional, *e.g.*, temperature sensitive.

Once validation of the inhibitor ORF is achieved, a bi-directional deletion analysis can be carried out using the same experimental system to identify the minimal polypeptide segment that has inhibitor activity. This may be carried out by a variety of means, *e.g.*, by exonuclease or PCR methodologies, and is used to determine if a relatively small segment of the ORF (*i.e.*, the product of the ORF) still possesses inhibitory activity when isolated away from its native sequence. If so, a portion of the ORF encoding this "active portion" can be used as a template for the synthesis of novel anti-microbial agents and further allowing derivation of the peptide sequence, *e.g.*, using modified peptides and/or peptidomimetics.

In creation of certain peptidomimetics, the peptide backbone is transformed into a carbon-based hydrophobic structure that can retain inhibitor activity against the bacterium. This is done by standard medicinal chemistry methods, typically monitored by measuring growth inhibition of the various molecules in liquid cultures or on solid medium. These mimetics can also represent lead compounds for the development of novel antibiotics.

Recently, a major effort has been undertaken by the pharmaceutical industry and their biotechnology partners for the sequencing of bacterial pathogen genomes. The rationale is that the systematic sequencing of the genome will identify all of the bacterial proteins and therefore this proteome will be the target for designing novel inhibitor antibiotics. Although systematic, this approach has several major problems. The first is that analysis of primary amino acid sequences of bacterial proteins does not immediately reveal which protein will be essential for viability of the bacterium, and target validation is thus a major issue. The second problem is one of redundancy, as several biochemical pathways are either structurally duplicated in bacteria (different isoforms of the same enzyme), or functionally duplicated by the presence of salvage pathways in the event of a metabolic block in one pathway (different nutritional conditions). The third is that even a valid target may not be structurally or

functionally amenable to inhibition by small molecules because of inaccessibility (sequestration of target).

Therefore, there is considerable interest within the pharmaceutical and biotechnology industry in identifying key targets for drug discovery amongst the mass
5 of novel targets generated by large-scale genomic sequencing projects.

On the other hand, and underscoring the instant invention, the phages herein described have, over millions of years, evolved specific mechanisms to target such key biochemical pathways and proteins. In the few cases where inhibition by phages has been elucidated (*e.g.*, see ref. 3), such bacterial targets are invariably rate-limiting
10 in their respective biochemical pathways, are not redundant, and/or are readily accessible for inhibition by the phage (or by another inhibitory compound). Therefore, the sixth step of this invention involves identifying the host biochemical pathways and proteins that are targeted by the phage inhibitory mechanisms.

15 Identifying, Validating, and Characterizing Bacterial Host Target Proteins and Affected Pathways

A rationale for this step is that the inhibitor ORF product from the phage physically interacts with and/or modifies certain microbial host components to block their function. Exemplary approaches which can be used to identify the host bacterial
20 pathways and proteins that interact with, and preferably also are inhibited by, phage ORF product(s) are described below.

One approach is a genetic screen to determine physiological protein:protein interaction, for example, using a yeast two hybrid system. In this assay, the phage ORF is fused to the carboxyl terminus of the yeast Gal4 activation domain II (amino
25 acids 768-881) to create a bait vector. A cDNA library of cloned *S. aureus* sequences which have been engineered into a plasmid where the *S. aureus* sequences are fused to the DNA binding domain of Gal4 is also generated. These plasmids are introduced alone, or in combination, into yeast strain Y190 - previously engineered with chromosomally integrated copies of the *E. coli lacZ* and the selectable HIS3 genes,
30 both under Gal4 regulation (Durfee, T., Becherer, K., Chen, P.-L., Yeh, S.-H., Yang, Y., Kilburn, A.E., Lee, W.-H., and Elledge, S.J. (1993). *Genes & Dev.* 7, 555-569). If the two proteins expressed in yeast interact, the resulting complex will activate transcription from promoters containing Gal4 binding sites. A *lacZ* and His3 gene, each driven by a promoter containing Gal4 binding sites, have been integrated into the
35 genome of the host yeast system used for measuring protein-protein interactions. Such a system provides a physiological environment in which to detect potential protein interactions. This system has been extensively used to identify novel protein-protein

interaction partners and to map the sites required for interaction (for example, to identify interacting partners of translation factors (Qiu, H., Garcia-Barrio, M.T., and Hinnebusch, A.G. (1998). *Mol & Cell Biology* 18, 2697-2711), transcription factors (Katagiri, T., Saito, H., Shinohara, A., Ogawa, H., Kamada, N., Nakamura, Y., and Miki, Y. (1998). *Genes, Chromosomes & Cancer* 21, 217-222), and proteins involved in signal transduction (Endo, T.A., Masuhara, M., Yokouchi, M., Suzuki, R., Sakamoto, H., Mitsui, K., Matsumoto, A., Tanimura, S., Ohtsubo, M., Misawa, H., Miyazaki, T., Leonor N., Taniguchi, T., Fujita, T., Kanakura, Y., Komiya, S., and Yoshimura, A. *Nature*. 387, 921-924). This approach has also been used in many published reports to identify interaction between mammalian viral and mammalian cell proteins.

For example, the non-structural protein NS1 of parvovirus is essential for viral DNA amplification and gene expression and is also the major cytopathic effector of these viruses. A yeast two-hybrid screen with NS1 identified a novel cellular protein of unknown function that interacts with NS-1, called SGT, for small glutamine-rich tetratricopeptide repeat (TPR)-containing protein (Cziepluch C. Kordes E. Poirey R. Grewenig A. Rommelaere, J. and Jauniaux JC. (1998) *J Virol.* 72, 4149-4156). In another screen, the adenovirus E3 protein was recently shown to interact with a novel tumor necrosis factor alpha-inducible protein and to modulate some of the activities of E3 (Li Y. Kang J. and Horwitz M.S. (1998). *Mol & Cell Biol.* 18, 1601-1610). In yet another recent screen, the herpes simplex virus 1 alpha regulatory protein ICP0 was found to interact with (and stabilize) the cell cycle regulator cyclin D3 (Kawaguchi Y. Van Sant C. and Roizman B. (1997). *J Virol.* 71, 7328-7336).

Another two-hybrid system for identifying protein:protein interactions is commercially available from STRATEGENE™ as the CYTO-TRAP™ system (Chang et al., *Strategies Newsletter* 11(3), 65-68 (1998)(from Stratagene)). The system is a yeast-based method for detecting protein:protein interactions *in vivo*, using activation of the Ras signal transduction cascade by localizing a signal pathway component, human Sos (hSos), to its activation site in the yeast plasma membrane. The system uses a temperature-sensitive *Saccharomyces cerevisiae* mutant, strain cdc25H, which contains a point mutation at amino acid residue 1328 of the cdc25 gene. This gene encodes a guanyl nucleotide exchange factor which binds and activates Ras, leading to cell growth. The mutation in the cdc25 gene prevents host growth at 37°C, but at a permissive temperature of 25°C, growth is normal. The system utilizes the ability of (hSos) to complement the cdc25 defect and activate the yeast Ras signaling pathway. Once (hSos) is expressed and localized to the plasma membrane, the cdc25H yeast strain grows at 37°C. Localizing hSos to the plasma

membrane occurs through a protein:protein interaction. A protein of interest, or bait, is expressed as a fusion protein with hSos. The library, or target proteins are expressed with the myristylation membrane-localization signal. The yeast cells are then incubated under restrictive conditions (37°C). If the bait and the target protein interact, the hSos protein is recruited to the membrane, activating the Ras signaling pathway and allowing the cdc25H yeast strain to grow at the restrictive temperature.

The protein targets of phage inhibitory ORFs can also be identified using bacterial genetic screens. One approach involves the overexpression of a phage inhibitory protein in mutagenized bacterial host species, followed by plating the cells and searching for colonies that can survive the antimicrobial activity of the inhibitory ORF. These colonies are then grown, their DNA extracted, and cloned into an expression vector that contains a replicon of a different incompatibility group from the plasmid expressing the original ORF. This library is then introduced into a wild-type host bacterium in conjunction with an expression vector driving synthesis of the phage ORF, followed by selection for surviving bacteria. Thus, bacterial DNA fragments from the survivors presumably contain a DNA fragment from the original mutagenized host bacterial genome that can protect the cell from the antimicrobial activity of the inhibitory phage ORF. This fragment can be sequenced and compared with that of the bacterial host to determine in which gene the mutation lies. This approach enables one to determine the targets and pathways that are affected by the killing function.

A second approach is based on identifying protein:protein interactions between the phage ORF product and bacterial *S. aureus*, e.g., proteins using a biochemical approach based, for example, on affinity chromatography. This approach has been used, for example, to identify interactions between lambda phage proteins and proteins from their *E. coli* host (Sopta, M., Carthew, R.W., and Greenblatt, J. (1985) *J. Biol. Chem.* 260, 10353-10369). The phage ORF is fused to a peptide tag (e.g. glutathione-S-transferase ("GST"), 6xHIS, ("HIS") and/or calmodulin binding protein ("CPB")) within a commercially available plasmid vector that directs high level expression on induction of a suitably responsive promoter driving the fusion's expression. The translated fusion protein is expressed in *E. coli*, purified, and immobilized on a solid phase matrix via, for example the tag. Total cell extracts from the host bacterium, e.g., *S. aureus*, are then passed through the affinity matrix containing the immobilized phage ORF fusion protein; host proteins retained on the column are then eluted under different conditions of ionic strength, pH, detergents etc., and characterized by gel electrophoresis and other techniques. Appropriate controls are run to guard against nonspecific binding to the resin. Target proteins thus

recovered should be enriched for the phage protein/peptide of interest and are subsequently electrophoretically or otherwise separated, purified, sequenced, or biochemically analyzed. Usually sequencing entails individual digestion of the proteins to completion with a protease (e.g.-trypsin), followed by molecular mass and amino acid composition and sequence determination using, for example, mass spectrometry, e.g., by MALDI-TOF technology (Qin, J., Fenyo, D., Zhao, Y., Hall, W.W., Chao, D.M., Wilson, C.J., Young, R.A. and Chait, B.T. (1997). *Anal. Chem.* 69, 3995-4001).

The sequence of the individual peptides from a single protein are then analyzed by the bioinformatics approach described above to identify the *S. aureus* protein interacting with the phage ORF. This analysis is performed by a computer search of the *S. aureus* genome for an identified sequence. Alternatively, all tryptic peptide fragments of the *S. aureus* genome can be predicted by computer software, and the molecular mass of such fragments compared to the molecular mass of the peptides obtained from each interacting protein eluted from the affinity matrix. The responsible gene sequence can be obtained, for example by using synthetic degenerate nucleic acid sequences to pull out the corresponding homologous bacterial sequence. Alternatively, antibodies can be generated against the peptide and used to isolate nascent peptide/mRNA transcript complexes, from which the mRNA can be reverse transcribed, cloned, and further characterized using the procedures discussed herein.

A variety of other binding assay methods are known in the art and can be used to identify interactions between phage proteins and bacterial proteins or other bacterial cell components. Such methods that allow or provide identification of the bacterial component can be used in this invention for identifying putative targets.

Validation of the interaction between the phage ORF product and the bacterial proteins or other components can be obtained by a second independent assay (e.g., co-immunoprecipitation or protein-protein crosslinking experiments (Qiu, H., Garcia-Barrio, M.T., and Hinnebusch, A.G. (1998). *Mol & Cell Biology* 18, 2697-2711; Brown, S. and Blumenthal, T. (1976). *Proc. Natl. Acad. Sci. USA* 73, 1131-1135)).

Finally, the essential nature of the identified bacterial proteins is preferably determined genetically by creating a constitutive or inducible partial or complete loss-of-function mutation in the gene encoding the identified interacting bacterial protein. This mutant is then tested for bacterial survival and replication.

The protein target of the phage inhibitor function can also be identified using a genetic approach. Two exemplary approaches will be delineated here. The first approach involves the overexpression of a predetermined phage inhibitor protein in mutagenized host bacteria, e.g., *S. aureus*, followed by plating the cells and searching

for colonies that can survive the inhibitor. These colonies will then be grown, their DNA extracted and cloned into an expression vector that contains a replicon of a different incompatibility group, and preferably having a different selectable marker than the plasmid expressing the phage inhibitor. Thus, host DNA fragments from the mutant that can protect the cell from phage ORF inhibition can be sequenced and compared with that of the bacterial host to determine in which gene the mutation lies. This approach allows rapid determination of the targets and pathways that are affected by the inhibitor.

Alternatively, the bacterial targets can be determined in the absence of selecting for mutations using an approach known as "multicopy suppression". In this approach, the DNA from the wild type host is cloned into an expression vector that can coexist, as previously described, with one containing a predetermined phage inhibitor. Those plasmids that contain host DNA fragments and genes that protect the host from the phage inhibitor can then be isolated and sequenced to identify putative targets and pathways in the host bacteria.

Regardless of the specific mode of identification, screening assays may additionally utilize gene fusions to specific "reporter genes" to identify a bacterial gene(s) whose expression is affected when the host target pathway is affected by the phage inhibitor. Such gene fusions can be used to search a number of small molecule compounds for inhibitors that may affect this pathway and thus cause cell inhibition. This approach will allow the screening of a large number of molecules on petri dishes or 96-well format by monitoring for a simple color change in the bacterial colonies. In this manner, we can validate host targets and classes of compounds for further study and clinical development. These inhibitors also represent lead compounds for the development of other antibiotics.

Bioinformatics and comparative genomics are preferably then applied to the identified bacterial gene products to predict biochemical function. The biochemical activity of the protein can be verified *in vitro* in cell free assays or *in vivo* in intact cells. *In vitro* biochemical assays utilizing cell-free extracts or purified protein are established as a basis for the screening and development of inhibitors.

These inhibitors, preferably small molecule inhibitors, may comprise peptides, antibodies, products from natural sources such as fungal or plant extracts or small molecule organic compounds. In general, small molecule organic compounds are preferred. These compounds may, for example, be identified within large compound libraries, including combinatorial libraries. For example, a plurality of compounds, preferably a large number of compounds can be screened to determine whether any of the compounds binds or otherwise disrupts or inhibits the identified bacterial target.

Compounds identified as having any of these activities can then be evaluated further in cell culture and/or animal model systems to determine the pharmacological properties of the compound, including the specific anti-microbial ability of the compound.

- 5 For mixtures of natural products, including crude preparations, once a preparation or fraction of a preparation is shown to have an anti-microbial activity, the active substance can be isolated and identified using techniques well known in the art, if the compound is not already available in a purified form.

- 10 Identified compounds possessing anti-microbial activity and similar compounds having structural similarity can be further evaluated and, if necessary, derivatized according to synthesis and/or modification methods available in the art selected as appropriate for the particular starting molecule.

Derivatization of identified anti-microbials

- 15 In cases where the identified anti-microbials above might represent peptidal compounds, the *in vivo* effectiveness of such compounds may be advantageously enhanced by chemical modification using the natural polypeptide as a starting point and incorporating changes that provide advantages for use, for example, increased stability to proteolytic degradation, reduced antigenicity, improved tissue penetration,
20 and/or improved delivery characteristics.

- In addition to active modifications and derivative creations, it can also be useful to provide inactive modifications or derivatives for use as negative controls or introduction of immunologic tolerance. For example, a biologically inactive derivative which has essentially the same epitopes as the corresponding natural
25 antimicrobial can be used to induce immunological tolerance in a patient being treated. The induction of tolerance can then allow uninterrupted treatment with the active anti-microbial to continue for a significantly longer period of time.

- Modified anti-microbial polypeptides and derivatives can be produced using a number of different types of modifications to the amino acid chain. Many such
30 methods are known to those skilled in the art. The changes can include, for example, reduction of the size of the molecule, and/or the modification of the amino acid sequence of the molecule. In addition, a variety of different chemical modifications of the naturally occurring polypeptide can be used, either with or without modifications to the amino acid sequence or size of the molecule. Such chemical modifications can,
35 for example, include the incorporation of modified or non-natural amino acids or non-amino acid moieties during synthesis of the peptide chain, or the post-synthesis modification of incorporated chain moieties.

The oligopeptides of this invention can be synthesized chemically or through an appropriate gene expression system. Synthetic peptides can include both naturally occurring amino acids and laboratory synthesized, modified amino acids.

Also provided herein are functional derivatives of anti-microbial proteins or polypeptides. By "functional derivative" is meant a "chemical derivative,"
5 "fragment," "variant," "chimera," or "hybrid" of the polypeptide or protein, which terms are defined below. A functional derivative retains at least a portion of the function of the protein, for example reactivity with a specific antibody, enzymatic activity or binding activity.

10 A "chemical derivative" of the complex contains additional chemical moieties not normally a part of the protein or peptide. Such moieties may improve the molecule's solubility, absorption, biological half-life, and the like. The moieties may alternatively decrease the toxicity of the molecule, eliminate or attenuate any undesirable side effect of the molecule, and the like. Moieties capable of mediating
15 such effects are disclosed in Alfonso and Gennaro (1995). Procedures for coupling such moieties to a molecule are well known in the art. Covalent modifications of the protein or peptides are included within the scope of this invention. Such modifications may be introduced into the molecule by reacting targeted amino acid residues of the peptide with an organic derivatizing agent that is capable of reacting
20 with selected side chains or terminal residues, as described below.

Cysteiny l residues most commonly are reacted with alpha-haloacetates (and corresponding amines), such as chloroacetic acid or chloroacetamide, to give carboxymethyl or carboxyamidomethyl derivatives. Cysteiny l residues also are derivatized by reaction with bromotrifluoroacetone, chloroacetyl phosphate, N-
25 alkylmaleimides, 3-nitro-2-pyridyl disulfide, methyl 2-pyridyl disulfide, p-chloro-mercuribenzoate, 2-chloromercuri-4-nitrophenol, or chloro-7-nitrobenzo-2-oxa-1,3-diazole.

Histidyl residues are derivatized by reaction with diethylprocarbonate at pH 5.5-7.0 because this agent is relatively specific for the histidyl side chain. Para-
30 bromophenacyl bromide also is useful; the reaction is preferably performed in 0.1 M sodium cacodylate at pH 6.0.

Lysiny l and amino terminal residues are reacted with succinic or other carboxylic acid anhydrides. Derivatization with these agents has the effect of reversing the charge of the lysiny l residues. Other suitable reagents for derivatizing
35 primary amine- containing residues include imidoesters such as methyl picolinimide; pyridoxal phosphate; pyridoxal; chloroborohydride;

trinitrobenzenesulfonic acid; O-methylisourea; 2,4 pentanedione; and transaminase-catalyzed reaction with glyoxylate.

Arginyl residues are modified by reaction with one or several conventional reagents, among them phenylglyoxal, 2,3-butanedione, 1,2-cyclohexanedione, and
5 ninhydrin. Derivatization of arginine residues requires that the reaction be performed in alkaline conditions because of the high pK_a of the guanidine functional group. Furthermore, these reagents may react with the groups of lysine as well as the arginine alpha-amino group.

Tyrosyl residues are well-known targets of modification for introduction of
10 spectral labels by reaction with aromatic diazonium compounds or tetranitromethane. Most commonly, N-acetylimidizol and tetranitromethane are used to form O-acetyl tyrosyl species and 3-nitro derivatives, respectively.

Carboxyl side groups (aspartyl or glutamyl) are selectively modified by
reaction carbodiimide ($R'-N-C-N-R'$) such as 1-cyclohexyl-3-(2-morpholinyl(4-ethyl)
15 carbodiimide or 1-ethyl-3-(4-azonia-4,4-dimethylpentyl) carbodiimide. Furthermore, aspartyl and glutamyl residues are converted to asparaginy and glutaminy residues by reaction with ammonium ions.

Glutaminy and asparaginy residues are frequently deamidated to the
corresponding glutamyl and aspartyl residues. Alternatively, these residues are
20 deamidated under mildly acidic conditions. Either form of these residues falls within the scope of this invention.

Derivatization with bifunctional agents is useful, for example, for cross-
linking component peptides to each other or the complex to a water-insoluble support
matrix or to other macromolecular carriers. Commonly used cross-linking agents
25 include, for example, 1,1-bis (diazoacetyl)-2-phenylethane, glutaraldehyde, N-hydroxysuccinimide esters, for example, esters with 4-azidosalicylic acid, homobifunctional imidoesters, including disuccinimidyl esters such as 3,3'-dithiobis(succinimidylpropionate), and bifunctional maleimides such as bis-N-maleimido-1,8-octane. Derivatizing agents such as methyl-3-[p-azidophenyl)
30 dithiolpropioimide yield photoactivatable intermediates that are capable of forming crosslinks in the presence of light. Alternatively, reactive water-insoluble matrices such as cyanogen bromide-activated carbohydrates and the reactive substrates described in U.S. Patent Nos. 3,969,287; 3,691,016; 4,195,128; 4,247,642; 4,229,537; and 4,330,440 are employed for protein immobilization.

35 Other modifications include hydroxylation of proline and lysine, phosphorylation of hydroxyl groups of seryl or threonyl residues, methylation of the alpha-amino groups of lysine, arginine, and histidine side chains (Creighton, T.E.,

Proteins: Structure and Molecular Properties, W.H. Freeman & Co., San Francisco, pp. 79-86 (1983)), acetylation of the N-terminal amine, and, in some instances, amidation of the C-terminal carboxyl groups.

Such derivatized moieties may improve the stability, solubility, absorption, biological half life, and the like. The moieties may alternatively eliminate or attenuate any undesirable side effect of the protein complex. Moieties capable of mediating such effects are disclosed, for example, in Alfonso and Gennaro (1995).

The term "fragment" is used to indicate a polypeptide derived from the amino acid sequence of the protein or polypeptide having a length less than the full-length polypeptide from which it has been derived. Such a fragment may, for example, be produced by proteolytic cleavage of the full-length protein. Preferably, the fragment is obtained recombinantly by appropriately modifying the DNA sequence encoding the proteins to delete one or more amino acids at one or more sites of the C-terminus, N-terminus, and/or within the native sequence.

Another functional derivative intended to be within the scope of the present invention is a "variant" polypeptide that either lacks one or more amino acids or contains additional or substituted amino acids relative to the native polypeptide. The variant may be derived from a naturally occurring polypeptide by appropriately modifying the protein DNA coding sequence to add, remove, and/or to modify codons for one or more amino acids at one or more sites of the C-terminus, N-terminus, and/or within the native sequence.

A functional derivative of a protein or polypeptide with deleted, inserted and/or substituted amino acid residues may be prepared using standard techniques well-known to those of ordinary skill in the art. For example, the modified components of the functional derivatives may be produced using site-directed mutagenesis techniques (as exemplified by Adelman et al., 1983, *DNA* 2:183; Sambrook et al., 1989) wherein nucleotides in the DNA coding sequence are modified such that a modified coding sequence is produced, and thereafter expressing this recombinant DNA in a prokaryotic or eukaryotic host cell, using techniques such as those described above. Alternatively, components of functional derivatives of complexes with amino acid deletions, insertions and/or substitutions may be conveniently prepared by direct chemical synthesis, using methods well-known in the art.

Insofar as other anti-microbial inhibitor compounds identified by the invention described herein may not be peptidal in nature, other chemical techniques exist to allow their suitable modification, as well, and according the desirable principles discussed above.

Administration and Pharmaceutical Compositions

For the therapeutic and prophylactic treatment of infection, the preferred method of preparation or administration of anti-microbial compounds will generally vary depending on the precise identity and nature of the anti-microbial being delivered. Thus, those skilled in the art will understand that administration methods known in the art will also be appropriate for the compounds of this invention.

The particularly desired anti-microbial can be administered to a patient either by itself, or in pharmaceutical compositions where it is mixed with suitable carriers or excipient(s). In treating an infection, a therapeutically effective amount of an agent or agents is administered. A therapeutically effective dose refers to that amount of the compound that results in amelioration of one or more symptoms of bacterial infection and/or a prolongation of patient survival or patient comfort.

Toxicity, therapeutic and prophylactic efficacy of anti-microbials can be determined by standard pharmaceutical procedures in cell cultures and/or experimental organisms such as animals, *e.g.*, for determining the LD₅₀ (the dose lethal to 50% of the population) and the ED₅₀ (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio LD₅₀/ED₅₀. Compounds that exhibit large therapeutic indices are preferred. The data obtained from these cell culture assays and animal studies can be used in formulating a range of dosage for use in humans. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED₅₀ with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized.

For any compound identified and used in the method of the invention, the therapeutically effective dose can be estimated initially from cell culture assays. Such information can be used to more accurately determine useful doses in organisms such as plants and animals, preferably mammals, and most preferably humans. Levels in plasma may be measured, for example, by HPLC or other means appropriate for detection of the particular compound.

The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition (see *e.g.* Fingl et. al., in *The Pharmacological Basis of Therapeutics*, 1975, Ch. 1 p.1).

It should be noted that the attending physician would know how and when to terminate, interrupt, or adjust administration due to toxicity, organ dysfunction, or other systemic malady. Conversely, the attending physician would also know to adjust treatment to higher levels if the clinical response were not adequate (precluding

toxicity). The magnitude of an administered dose in the management of the disorder of interest will vary with the severity of the condition to be treated and the route of administration. The severity of the condition may, for example, be evaluated, in part, by standard prognostic evaluation methods. Further, the dose and perhaps dose
5 frequency, will also vary according to the age, body weight, and response of the individual patient. A program comparable to that discussed above also may be used in veterinary or phyto medicine.

Depending on the specific infection target being treated and the method selected, such agents may be formulated and administered systemically or locally, i.e.,
10 topically. Techniques for formulation and administration may be found in Alfonso and Gennaro (1995). Suitable routes may include, for example, oral, rectal, transdermal, vaginal, transmucosal, intestinal, parenteral, intramuscular, subcutaneous, or intramedullary injections, as well as intrathecal, intravenous, or intraperitoneal injections.

15 For injection, the agents of the invention may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks' solution, Ringer's solution, or physiological saline buffer. For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

20 Use of pharmaceutically acceptable carriers to formulate identified anti-microbials of the present invention into dosages suitable for systemic administration is within the scope of the invention. With proper choice of carrier and suitable manufacturing practice, the compositions of the present invention, in particular those formulated as solutions, may be administered parenterally, such as by intravenous
25 injection. Appropriate compounds can be formulated readily using pharmaceutically acceptable carriers well known in the art into dosages suitable for oral administration. Such carriers enable the compounds of the invention to be formulated as tablets, pills, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated.

30 Agents intended to be administered intracellularly may be administered using techniques well known to those of ordinary skill in the art. For example, such agents may be encapsulated into liposomes, then administered as described above. Liposomes are spherical lipid bilayers with aqueous interiors. All molecules present in an aqueous solution at the time of liposome formation are incorporated into the
35 aqueous interior. The liposomal contents are both protected from the external microenvironment and, because liposomes fuse with cell membranes, are efficiently

delivered into the cell cytoplasm. Additionally, due to their hydrophobicity, small organic molecules may be directly administered intracellularly.

Pharmaceutical compositions suitable for use in the present invention include compositions wherein the active ingredients are contained in an effective amount to
5 achieve the intended purpose. Determination of the effective amounts is well within the capability of those skilled in the art.

In addition to the active ingredients, these pharmaceutical compositions may contain suitable pharmaceutically acceptable carriers comprising excipients and auxiliaries which facilitate processing of the active compounds into preparations
10 which can be used pharmaceutically. The preparations formulated for oral administration may be in the form of tablets, dragees, capsules, or solutions, including those formulated for delayed release or only to be released when the pharmaceutical reaches the small or large intestine.

The pharmaceutical compositions of the present invention may be
15 manufactured in a manner that is itself known, *e.g.*, by means of conventional mixing, dissolving, granulating, dragee-making, levitating, emulsifying, encapsulating, entrapping or lyophilizing processes.

Pharmaceutical formulations for parenteral administration include aqueous solutions of the active anti-microbial compounds in water-soluble form.
20 Alternatively, suspensions of the active compounds may be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acid esters, such as ethyl oleate or triglycerides, or liposomes. Aqueous injection suspensions may contain substances which increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or
25 dextran. Optionally, the suspension may also contain suitable stabilizers or agents which increase the solubility of the compounds to allow for the preparation of highly concentrated solutions.

Pharmaceutical preparations for oral use can be obtained by combining the active compounds with solid excipient, optionally grinding a resulting mixture, and
30 processing the mixture of granules, after adding suitable auxiliaries, if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium
35 carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate.

Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent mixtures.

- 5 Dyestuffs or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

- Pharmaceutical preparations which can be used orally include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active ingredients in admixture with filler such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition, stabilizers may be added.

- 15 The above methodologies may be employed either actively or prophylactically against an infection of interest.

Computer-related Aspects and Embodiments

- In addition to the provision of compounds as chemical entities, nucleotide sequences, or fragments thereof at least 95%, preferably at least 97%, more preferably at least 99%, and most preferably at least 99.9% identical to phage inhibitor sequences can also be provided in a variety of additional media to facilitate various uses.

- Thus, as used in this section, "provided" refers to an article of manufacture, rather than an actual nucleic acid molecule, which contains a nucleotide sequence of the present invention; *e.g.*, a nucleotide sequence of an exemplary bacteriophage or a sequence encoding a bacterial target or a fragment thereof, preferably a nucleotide sequence at least 95%, more preferably at least 99% and most preferably at least 99.9% identical to such a bacteriophage or bacterial sequence, for example, to a polynucleotide of an unsequenced phage listed in Table 1, preferably of bacteriophage 77 (*S. aureus* host) or bacteriophage 3A (*S. aureus* host) or bacteriophage 96 (*S. aureus* host). Such an article provides a large portion of the particular bacteriophage genome or bacterial gene and parts thereof (*e.g.*, a bacteriophage open reading frame (ORF)) in a form which allows a skilled artisan to examine and/or analyze the sequence using means not directly applicable to examining the actual genome or gene, or subset thereof as it exists in nature or in purified form as a chemical entity.

In one application of this aspect, a nucleotide sequence of the present invention can be recorded on computer readable media. As used herein, "computer

readable media" refers to any medium that can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories, such as magnetic/optical storage media. A skilled artisan can readily appreciate how any of the presently known computer readable mediums can be used to create an article of manufacture which includes one or more computer readable media having recorded thereon a nucleotide sequence or sequences of the present invention. Likewise, it will be clear to those of skill how additional computer readable media that may be developed also can be used to create analogous manufactures having recorded thereon a nucleotide sequence of the present invention.

As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the presently known methods for recording information on computer readable medium to generate manufactures comprising the nucleotide sequence information of the present invention.

A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of data processor programs and formats can be used to store the nucleotide sequence information of the present invention on computer readable medium. The sequence information can, for example, be presented in a word processing text file, formatted in commercially available software such as WordPerfect and Microsoft Word, or represented in the form of an ASCII file, stored in a database application, such as DB2, Sybase, Oracle, or the like. A skilled artisan can readily adapt any number of data processor structuring formats (e.g., text file or database) in order to obtain computer readable medium having recorded thereon the nucleotide sequence information of the present invention.

Computer software is publicly available which allows a skilled artisan to access sequence information provided in a computer readable medium. Thus, by providing in computer readable form a nucleotide sequence of an unsequenced bacteriophage, such as an exemplary bacteriophage listed in Table 1 or of a sequence encoding a bacterial target or a fragment thereof, preferably a nucleotide sequence at least 95%, more preferably at least 99% and most preferably at least 99.9% identical to such a bacteriophage or bacterial sequence, for example, to a polynucleotide of bacteriophage 77 (*S. aureus* host) or bacteriophage 3A (*S.aureus* host) bacteriophage

96 (*S. aureus* host), bacteriophage 44AHJD (*S. aureus* host), bacteriophage Dp-1 (*Streptococcus pneumoniae* host), or bacteriophage 182 (*Enterococcus* host) the present invention enables the skilled artisan to routinely access the provided sequence information for a wide variety of purposes.

5 Those skilled in the art understand that software can implement a variety of different search or analysis software which implement sequence search and analysis algorithms, *e.g.*, the BLAST (Altschul et al., J. Mol. Biol. 215:403-410 (1990) and BLAZE (Brutlag et al., Comp. Chem 17:203-207 (1993)) search algorithms. For example, such search algorithms can be implemented on a Sybase system and used to
10 identify open reading frames (ORFs) within the bacteriophage genome which contain homology to ORFs or proteins from other viruses, *e.g.*, other bacteriophage, and other organisms, *e.g.*, the host bacterium. Among the ORFs discussed herein are protein encoding fragments of the bacteriophage genomes which encode bacteria-inhibiting proteins or fragments.

15 The present invention further provides systems, particularly computer-based systems, which contain the sequence information described. Such systems are designed to identify, among other things, useful fragments of the bacteriophage genomes.

 As used herein, "a computer-based system" refers to the hardware, software,
20 and data storage media used to analyze the nucleotide sequence information of the present invention. The minimum hardware of the computer-based systems of the present invention comprises a central processing unit (CPU), input device, output device, and data storage medium or media. A skilled artisan will readily recognize that any of the currently available general purpose computer-based system are suitable
25 for use in the present invention, as well as a variety of different specialized or dedicated computer-based systems.

 As stated above, the computer-based systems of the present invention comprise data storage media having stored therein a nucleotide sequence of the present invention and the necessary hardware and software for supporting and
30 implementing a search and/or analysis program.

 As used herein, "data storage media" refers to memory which can store nucleotide sequence information of the present invention, or a memory access means which can access manufactures having recorded thereon the nucleotide sequence information of the present invention.

35 As used herein, "search program" refers to one or more programs which are implemented on the computer-based system to compare a target sequence or target structural motif with the sequence information stored within the data storage means.

Search means are used to identify fragments or regions of the present genomic sequences which match a particular target sequence or target motif. A variety of known algorithms are disclosed publicly and a variety of commercially available software for conducting search means are and can be used in the computer-based systems of the present invention. Examples of such software includes, but is not limited to, MacPattern (EMBL), BLASTN and BLASTX (NCBIA). A skilled artisan can readily recognize that any one of the available algorithms or implementing software packages for conducting homology searches and/or sequence analyses can be adapted for use in the present computer-based systems.

As used herein in connection with sequence searches and analyses, a "target sequence" can be any DNA or amino acid sequence of six or more nucleotides or two or more amino acids. A skilled artisan can readily recognize that the longer a target sequence is, the less likely a target sequence will be present as a random occurrence in the database. Also, the target sequence length is preferably selected to include sequence corresponding to a biologically relevant portion of an encoded product, for example a region which is expected to be conserved across a range of source organisms. Preferably the sequence length of a target polypeptide sequence is from 5-100 amino acids, more preferably 7-50 or 7-100 amino acids, and still more preferably 10-80 or 10-100 amino acids. Preferably the sequence length of a target polynucleotide sequence is from 15-300 nucleotide residues, more preferably from 21-240 or 21-300, and still more preferably 30-150 or 30-300 nucleotide residues. However, it is well recognized that searches for commercially important fragments, such as sequence fragments involved in gene expression and protein processing, may be of shorter length. Likewise, it may be desirable to search and/or analyze longer sequences.

As used herein, "a target structural motif," or "target motif," refers to any rationally selected sequence or combination of sequences in which the sequence(s) are chosen based on a three-dimensional configuration which is formed upon the folding of the target motif. There are a variety of target motifs known in the art. Protein target motifs include, but are not limited to, enzymatic active sites and signal sequences. Nucleic acid target motifs include, but are not limited to promoter sequences, hairpin structures and inducible expression elements (protein binding sequences).

A variety of structural formats for the input and output devices can be used to input and output the information in the computer-based systems of the present invention. A preferred format for an output device ranks fragments of the bacteriophage or bacterial sequences possessing varying degrees of homology to the

target sequence or target motif. Such presentation provides a skilled artisan with a ranking of sequences which contain various amounts of the target sequence or target motif and identifies the degree of homology contained in the identified fragment.

A variety of comparing methods and/or devices and/or formats can be used to
5 compare a target sequence or target motif with the sequence stored in data storage media to identify sequence fragments of the bacteriophage or bacterium in question. One skilled in the art can readily recognize that any one of the publicly available homology search programs can be used as the search program for the computer-based systems of the present invention. Of course, suitable proprietary systems that may be
10 known to those of skill, or later developed, also may be employed in this regard.

Figure 6 provides a block diagram of a computer system illustrative of embodiments of this aspect of present invention. The computer system 102 includes a processor 106 connected to a bus 104. Also connected to the bus 104 are a main memory 108 (preferably implemented as random access memory, RAM) and a variety
15 of secondary storage devices 110, such as a hard drive 112 and a removable medium storage device 114. The removable medium storage device 114 may represent, for example, a floppy disk drive, a CD-ROM drive, a magnetic tape drive, etc. A removable storage medium 116 (such as a floppy disk, a compact disk, a magnetic tape, etc.) containing control logic and/or data recorded therein may be inserted into
20 the removable medium storage device 114. The computer system 102 includes appropriate software for reading the control logic and/or the data from the removable medium storage device 114, once it is inserted into the removable medium storage device 114.

A nucleotide sequence of the present invention may be stored in a well-known
25 manner in the main memory 108, any of the secondary storage devices 110, and/or a removable storage medium 116. During execution, software for accessing and processing the sequence (such as search tools, comparing tools, etc.) reside in main memory 108, in accordance with the requirements and operating parameters of the operating system, the hardware system and the software program or programs.

30 The data storage medium in which the sequence is embodied and the central processor need not be part of a single stand-alone computer, but may be separated so long as data transfer can occur. For example, the processor or processors being utilized for a search or analysis can be part of one general purpose computer, and the data storage medium can be part of a second general purpose computer connected to a
35 network, or the data storage medium can be part of a network server. As another example the data storage medium can be part of a computer system or network accessible over telephone lines or other remote connection method.

EXAMPLES

Example 1. Growth of *Staph A* bacteriophage 77 and purification of genomic DNA.

5 The *Staphylococcus aureus* propagating strain (PS 77; ATCC #27699) was used as a host to propagate its respective phage 77 (ATCC # 27699-B1). Two rounds of plaque purification of phage 77 were performed on soft agar essentially as described in Sambrook et al (1989). Briefly, the PS 77 strain was grown overnight at 37°C in Nutrient broth [NB: 0.3% Bacto beef extract, 0.5% Bacto peptone (Difco
10 Laboratories) and 0.5% NaCl (w/v)]. The culture was then diluted 20x in NB and incubated at 37°C until the $OD_{540} = .2$ (early log phase) with constant agitation. In order to obtain single plaques, phage 77 was subjected to 10-fold serial dilutions using phage buffer (1 mM $MgSO_4$, 5 mM $MgCl_2$, 80 mM NaCl and 0.1% Gelatin (w/v)) and 10 μ l of each dilution was used to infect 0.5 ml of the cell suspension in the presence
15 of 400 μ g/ml $CaCl_2$. After incubation of 15 min at room temperature (RT), 2 ml of melted soft agar kept at 45°C (NB supplemented with 0.6% agar) was added to the mixture and poured onto the surface of 100 mm nutrient agar plates (0.3% Bacto Beef extract, 0.5% Bacto peptone, 0.5% NaCl and 1.5% Bacto agar (w/v)). After overnight incubation at 30°C, a single plaque was isolated, resuspended in 1 ml of phage buffer
20 by end over end rotation for 2 hrs at 20°C, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 30°C, a single plaque was isolated and used as a stock.

 The propagation procedure for bacteriophage 77 was modified from the agar layer method of Swanstörn and Adams (1951). Briefly, the PS 77 strain was grown to
25 stationary phase overnight at 37°C in Nutrient broth. The culture was then diluted twenty-fold in NB and incubated at 37°C until the $OD_{540} = .2$. The suspension (15×10^7 Bacteria) was then mixed with 15×10^5 plaque forming units (pfu) to give a ratio of 100-bacteria/phage particle in the presence of 400 μ g/ml of $CaCl_2$. After incubation for 15 min at 20°C, 7.5 ml of melted soft agar (NB plus 0.6% agar) were added to the
30 mixture and poured onto the surface of 150 mm nutrient agar plates and incubated 16 hrs at 30°C. To collect the phage plate lysate, 20 ml of NB were added to each plate and the soft agar layer was collected by scrapping off with a clean microscope slide followed by shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 RPM (2,830xg) in a JA-10 rotor
35 (Beckman) and the supernatant fluid (lysate) was collected and subjected to a treatment with 10 μ g /ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) PEG 8000 and

0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO₄, 5 mM MgCl₂, 80 mM NaCl and 0.1% Gelatin). The phage suspension was
5 extracted with 1 volume of chloroform and further purified by centrifugation on a cesium chloride step gradient as described in Sambrook et al. (1989), using a TLS 55 rotor centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 h at 28,000 rpm (67,000xg) at 4°C. Banded phage was collected and ultracentrifuged again on an isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000xg) for 24 h at
10 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 h at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl₂. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 mg/ml Proteinase K and 0.5% SDS and incubating for 1 h at 65°C, followed by successive extractions with 1 volume of
15 phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris pH 8.0, 1mM EDTA).

Example 2. DNA sequencing of Bacteriophage 77 genome

Four micrograms of phage 77 DNA was diluted in 200 µl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed
20 (550 Sonic Dismembrator™, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4 cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0])
25 as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 µl of 1 mM Tris (pH 8.5).

The ends of the sonicated DNA fragments were repaired with a combination of
30 T4 DNA polymerase and the Klenow fragment of E. coli DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume, 100 µl) containing sonicated phage DNA, 10 mM Tris-HCl [pH 8.0], 50 mM NaCl, 10 mM MgCl₂, 1 mM DTT, 50 µg/ml BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5
35 units of Klenow large fragment (New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the

DNA was precipitated with ethanol and the final DNA pellet was resuspended in 20 µl of H₂O.

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs)-treated pKS II+ vector (Stratagene). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 µl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 µl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10β according to standard procedures (Sambrook et al., 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 µl LB and 100 µg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *Hinc* II cloning site of the pKS II+ vector. PCR amplification of foreign insert was performed in a 15 µl reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl₂, 0.02% gelatin, 1 µM primer, 187.5 µM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 57°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using QIAprep™ spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye™ primer or ABI prism Big Dye™ terminator cycle sequencing ready reaction kit (Applied Biosystems). To ensure co-linearity of the sequence data and the genome, all regions of phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism Big Dye™ terminator cycle sequencing ready reaction kit.

Example 3. Bioinformatic management of primary nucleotide sequence from Phage 77.

Phage 77 sequence contigs were assembled using Sequencher™ 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of

the contigs. Phage DNA was used directly as sequencing template employing ABI prism BIG DYE™ terminator cycle sequencing ready reaction kit. The complete sequence of bacteriophage 77 is shown in Table 2.

A software program was developed and used on the assembled sequence of bacteriophage 77 to identify all putative ORFs larger than 33 codons. Other ORF identification software can also be utilized, preferably programs which allow alternative start codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI (<http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c>) for the bacterial genetic code.

When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons (start and stop codons) is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence.

Sequence homology (BLAST) searches for each ORF are then carried out using an implementation of BLAST programs, although any of a variety of different sequence comparison and matching programs can be utilized as known to those skilled in the art. Downloaded public databases used for sequence analysis include:

- i) non-redundant GenBank (<ftp://ncbi.nlm.nih.gov/blast/db/nr.Z>),
- ii) Swissprot (<ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z>);
- iii) vector (<ftp://ncbi.nlm.nih.gov/blast/db/vector.Z>);
- iv) pdbaa databases (<ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z>);
- v) *S. aureus* NCTC 8325 (<ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa>);
- vi) streptococcus pyogenes (<ftp://ftp.genome.ou.edu/pub/strep/strep-1k.fa>);
- vii) *Streptococcus pneumoniae* (ftp://ftp.tigr.org/pub/data/s_pneumoniae/gsp.contigs.112197.Z);
- viii) *Mycobacterium tuberculosis* CSU#9 (ftp://ftp.tigr.org/pub/data/m_tuberculosis/TB_091097.Z) and
- ix) pseudomonas aeruginosa (<http://www.genome.washington.edu/pseudo/data.html>).

The results of the homology searches performed on the ORFs is shown in Table 5.

Example 4. Subcloning of Bacteriophage 77 ORFs into a Staph A inducible expression system.

The shuttle vector pT0021, in which the firefly luciferase (*lucFF*) expression is controlled by the *ars* (arsenite) promoter/operator (Tauriainen et al., 1997), was modified in the following fashion. Two oligonucleotides corresponding to a short antigenic peptide derived from the hemagglutinin protein of influenza virus (HA epitope tag) were synthesized (Field et al., 1988). The sense strand HA tag sequence (with *Bam*HI, *Sal*I and *Hind*III cloning sites) is:

5'-gatcccggtcgaccaagctTACCCATACGACGTCCCAGACTACGCCAGCTGA-3'

(where upper case letters denote the nucleotide sequence of the HA tag); the antisense strand HA tag sequence (with a *Hind*III cloning site) is:

5'-agctTCAGCTGGCGTAGTCTGGGACGTCGTATGGGTAAagcttggtcgaccgg-3'

(where upper case letters denote the sequence of the HA tag). The two HA tag oligonucleotides were annealed and ligated into pT0021 vector which had been digested with *Bam*HI and *Hind*III. This manipulation resulted in replacement of the *lucFF* gene by the HA tag. This modified shuttle vector containing the *arsenite* inducible promoter, the *arsR* gene, and HA tag was named pTHA. A diagram outlining our modification of pT0021 to generate pTHA is shown in Fig. 1A.

Each ORF, encoded by Bacteriophage 77, larger than 33 amino acids and having a Shine-Dalgarno sequence upstream of the initiation codon was selected for functional analysis for bacterial inhibition. In total, 98 ORFs were selected and screened as detailed below. A list of these is presented in Table 3. Each individual ORF, from initiation codon to last codon (excluding the stop codon), was amplified from phage genomic DNA using the polymerase chain reaction (PCR). For PCR amplification of ORFs, each sense strand primer targets the initiation codon and is preceded by a *Bam*HI restriction site (5'-cgggatcc-3') and each antisense oligonucleotide targets the penultimate codon (the one before the stop codon) of the ORF and is preceded by a *Sal*I restriction site (5'-gagtcgaccg-3'). The PCR product of each ORF was gel purified and digested with *Bam*HI and *Sal*I. The digested PCR product was then gel purified using the Qiagen kit as described, ligated into *Bam*HI and *Sal*I digested pTHA vector, and used to transform *E. coli* bacterial strain DH10 β (as described above). As a result of this manipulation, the HA tag is set inframe with the ORF and is positioned at the carboxy terminus of each ORF (pTHA/ORF clones). Recombinant pTHA/ORF clones were picked and their insert sizes were confirmed by PCR analysis

using primers flanking the cloning site. The names and sequences of the primers that were used for the PCR amplification were: HAF:

5' TATTATCCAAACTTGAACA 3'; HAR: 5' CGGTGGTATATCCAGTGATT 3'. The sequence integrity of cloned ORFs was verified directly by DNA sequencing using primers HAF and HAR. In cases where verification of ORF sequence could not be achieved by one pass with the sequencing primers, additional internal primers were selected and used for sequencing.

Staphylococcus aureus strain RN4220 (Kreiswirth et al., 1983) was used as a recipient for the expression of recombinant plasmids. Electoporation was performed essentially as previously described (Schenk and Laddaga, 1992). Selection of recombinant clones was performed on Luria-Broth agar (LB-agar) plates containing 30 µg/ml of kanamycin.

For each ORF introduced in the pTHA plasmid, 3 independent transformants were isolated and used to individually inoculate cultures in 5 ml of TSB containing 30 µg/ml kanamycin, followed by growth to saturation (16 hrs at 30°C). An aliquot of this stationary phase culture was used to generate a frozen glycerol stock of the transformant (stored at -80°C). The remaining culture was used for plasmid DNA extraction. Bacterial cells were harvested by centrifugation at 3000 x g at 22°C for 5 min. The pellet was resuspended in 200 µl 25% sucrose containing 25U/ml of lysostaphin and incubated for 15 min at 37°C. Then, 400 µl of alkaline SDS solution (3% SDS, 0.2N NaOH) were added, well mixed and incubated for 7 min at room temperature. After the alkaline SDS treatment, 300 µl of ice-cold 3M sodium acetate pH 4.8 were added, and the mix is immediately spun at 13000g for 15 min at room temperature. The supernatant was transferred to a new 1.5 ml conical centrifuge tube and 650 µl of isopropanol (stored at room temperature) were added. The mix was then centrifuged at 13,000 x g for 5 min. The supernatant fluid was discarded, the pellet washed with 70% ethanol, and resuspended in 320 µl sterile distilled water.

The presence of individual phage 77 ORF DNA inserts in the plasmid was verified by PCR amplification using 1.5 µl transformant miniprep DNA in a PCR with primers flanking the cloning site of ORF in pTHA vector (HAF and HAR). The composition of the PCR reaction and the cycling parameters are identical to those employed for library screening described above.

Example 5. Functional assay for bacterial inhibitory activity of bacteriophage 77 ORFs.

The anti-microbial activity of individual phage 77 ORFs was monitored by two growth inhibitory assays, one on solid agar medium, the other in liquid medium.

In general, *Staphylococcus* bacteria transformed with expression plasmids containing individual ORFs were grown in normal TSA medium and stored in 19% glycerol. At pre-determined times, arsenite was added to the culture to induce transcription of the phage 77 ORFs cloned immediately downstream from an arsenite-inducible promoter in the pTHA expression plasmid.

The effect of ORF induction on bacterial growth characteristics was then monitored and quantitated. The growth inhibition assay on solid medium was performed by streaking pTHA/ORF containing *S. aureus* transformant onto LB-Kn and TSA-Kn plates containing increasing concentrations of sodium arsenite (0; 2.5; 5; and 7.5 μ M). Arsenite is used to induce the expression of cloned DNA in pTHA vector. In parallel, 3 μ l of 1/10 and 1/100 dilutions of the frozen cultures of the pTHA/ORF transformants were spotted as single drops onto LB-Kn and TSA-Kn plates containing increasing concentration of sodium arsenite (0; 2.5; 5; and 7.5 μ M). The plates were then incubated 16 hrs at 37°C, and the effect of arsenite-induced ORF expression on bacterial growth was monitored and quantitated by comparing the extent to that seen in control plates. As positive controls for growth inhibition, the *holin/lysin* genes of the *Staphylococcus aureus* phage Twort (Loessner et al., 1998) was subcloned into the pTHA *ars* inducible vector and used.

For the growth inhibition assay in liquid medium, stationary phase cultures were prepared by inoculating 2.5ml TSB-Kn with frozen *S. aureus* RN4220 transformants containing phage 77 ORFs cloned in pTHA vector followed by incubation for 16 hrs at 37°C. These cultures were then diluted 1/100 in the same medium, and the bacteria were allowed to grow for 2 hrs at 37°C to reach early log phase. 150 μ l of such culture were then mixed with 2.35 ml TSB-Kn medium with or without arsenite (the final concentration of arsenite in the medium was 0 or 5 μ M arsenite). After 3.5 hrs incubation at 37°C with shaking at 250 rpm, 100 μ l of bacterial culture was removed from each tube for OD₅₆₅ measurement. Serial ten-fold dilutions of the culture in buffered saline solution (0.85% NaCl) were then spotted onto TSB-Kn plates. The plates were incubated at 37°C 16 hrs and the number of surviving colonies counted the following day. The growth inhibitory property of individual ORFs was then quantitated by comparing CFU numbers under normal or arsenite-induction conditions. A schematic flow of the inhibition analysis is shown in Fig. 3 (also applicable to inhibition analysis for the other phage and bacteria pointed out herein). Inhibition results are shown in Figures 4A-C.

Example 6: Identification of Cecropin Signature Motif in *Staphylococcus aureus* Bacteriophage 3A ORF

The genome for *S. aureus* bacteriophage 3A was determined and the sequence was analyzed essentially as described for bacteriophage 77 in the examples above. Upon blast analysis of the identified open reading frames of phage 3A, the presence of an amino acid sequence corresponding to a cecropin signature motif was observed.

- 5 This motif (WDGHKTLEK) is located at position aa 481-489. Cecropins were originally identified in proteins from the cecropia moth and are recognized as potent antibacterial proteins that constitute an important part of the cell-free immunity of insects. Cecropins are small proteins (31-39 amino acid residues) that are active against both Gram-positive and Gram-negative bacteria by disrupting the bacterial
10 membranes. Although the mechanisms by which the cecropins cause cell death are not fully understood, it is generally thought to involve channel formation and membrane destabilization.

- The identification of a motif corresponding to a known inhibitor suggests that the product of ORF002 is also an inhibitory compound. Such inhibitory activity can
15 be confirmed as described herein or by other methods known in the art. Confirmation of the inhibitory activity would indicate that the ORF product could serve as the basis for construction of mimetic compounds and other inhibitors directed to the target of the ORF002 product.

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20 Boman, 1991, *Cell* 65:205-207.
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Example 7. Growth of *Staphylococcus aureus* bacteriophage 44AHJD:

- 25 *Staphylococcus aureus* propagating strain (PS 44A) (Felix d'Herelle Reference Centre #HER 1101) was used as a host to propagate its respective phage 44AHJD (Felix d'Herelle Reference Centre #HER 101). Two rounds of plaque purification of phage 44AHJD were performed on soft agar essentially as described in Sambrook *et al.* (1989). Briefly, the *Staphylococcus aureus* PS strain was grown overnight at 37°C
30 in Nutrient Broth [NB: 3 g Bacto Beef Extract, 5 g Bactopeptone per liter, (Difco Laboratories # 0003-17-8), supplemented with 0.5% NaCl]. The culture was then diluted 20 fold in NB and incubated at 37°C until an OD₅₄₀ of 0.2. In order to obtain single plaques, phage 44AHJD was subjected to 10-fold serial dilutions using the phage buffer (1 mM MgSO₄, 5 mM MgCl₂, 80 mM NaCl and 0.1% Gelatin) and 10 µl
35 were used to infect 0.5 ml of the cell suspension in the presence of 400 µg/ml of

CaCl₂. After incubation of 15 min at room temperature, 2 ml of melted soft agar (NB supplemented with 0.6% of agar) were added to the mixture and poured onto the surface of 100 mm nutrient agar plates (3 g Bacto Beef extract, 5 g Bactopeptone, 0.5% NaCl and 15 g of Bacto agar per liter (Difco Laboratories # 0001-17-0). After
5 overnight incubation at 37°C, a single plaque was isolated, resuspended in 1ml of phage buffer by end over end rotation for 2 h at room temperature and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock.

Large scale purification of bacteriophage and preparation of phage DNA was
10 as follows.

The propagation method was carried out by using the agar layer method described by Swanstörn and Adams (1951). Briefly, the PS 44A strain was grown to stationary phase overnight at 37°C in Nutrient Broth. The culture was then diluted 20x in NB and incubated at 37°C until the A₅₄₀ = 0.2. The suspension (15x10⁷ Bacteria)
15 was then mixed with 15x10⁵ phage particles to give a ratio of 100-bacteria/phage particle in the presence of 400 µg/ml of CaCl₂. After incubation of 15 min at room temperature, 7.5 ml of melted soft agar were added to the mixture and poured onto the surface of 150 mm nutrient agar plates and incubated overnight at 37°C. To collect the lysate, 20 ml of NB were added to each plate and the soft agar layer was collected by
20 scrapping off with a clean microscope slide and shaken vigorously for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant (lysate) is collected and subjected to a treatment with 10 µg/ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, 10% (w/v) of PEG 8000 and 0.5 M of NaCl were
25 added to the lysate and the mixture was incubated on ice for 16 h. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman).

The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO₄, 5 mM MgCl₂, 80 mM NaCl and 0.1% Gelatin). The phage suspension was extracted with 1
30 volume of chloroform and further purified by centrifugation on a preformed cesium chloride step gradient as described in Sambrook *et al.* (1989), using a TLS 55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 h at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an

isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 x g) for 24 h at 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 h at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl₂. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 µg/ml Proteinase K and 0.5% SDS and incubating for 1 h at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

Example 8. DNA sequencing of the Bacteriophage 44 AHJD genome.

Four mg of phage DNA was diluted in 200 µl of TE pH 8.0 in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4 cycles and size fractionated on 1% agarose gels. The sonicated DNA was then size fractionated by gel electrophoresis. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen) and eluted in 50 µl of 1mM Tris-HCl [pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of *E. coli* DNA polymerase 1 as follows. Reactions were performed in a final volume of 100 µl containing DNA, 10 mM Tris-HCl pH 8.0, 50 mM NaCl, 10 mM MgCl₂, 1 mM DTT, 5 µg BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of Klenow fragment (New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was ethanol precipitated and resuspended in 20 µl of H₂O.

Cloning of the sonicated phage DNA into pKSII vector and transformation:

Blunt-ended DNA fragments were cloned by ligation directly into the *HincII* site of the pKSII vector (Stratagene) dephosphorylated with calf intestinal alkaline phosphatase (New England Biolabs). A typical reaction contained 100 ng of vector, 2

to 5 µl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 µl containing 800 units of T4 DNA ligase (New England Biolabs) overnight at 16°C. Transformation and selection of positive clones was performed in the host strain DH10 β of *E. coli* using ampicillin as a selective antibiotic as described in Sambrook *et al.* (1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 µl LB and 100 µg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *HincII* cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 µl reaction volume containing 10 mM Tris-HCl (pH 8.3), 50 mM KCl, 1.5 mM MgCl₂, 0.02% gelatin, 1 mM primer, 187.5 µM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 58°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep™ spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism BigDye™ primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism BigDye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism BigDye™ terminator cycle sequencing ready reaction kit.

Example 9. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher™ 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI

prism BigDye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Staphylococcus aureus* bacteriophage 44AHJD is shown in Table 16.

A software program was used on the assembled sequence of bacteriophage 44AHJD to identify all putative ORFs larger than 33 codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI(<http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c>) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence. The predicted ORFs for bacteriophage 44AHJD are listed in Tables 17 & 18.

Sequence homology searches for each ORF were carried out using an implementation of blast programs. Downloaded public databases used for sequence analysis include:

- (i) non-redundant GenBank (<ftp://ncbi.nlm.nih.gov/blast/db/nr.Z>),
- ii) Swissprot (<ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z>);
- 25 iii) vector (<ftp://ncbi.nlm.nih.gov/blast/db/vector.Z>);
- iv) pdbaa databases (<ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z>);
- v) *Staphylococcus aureus* NCTC 8325 (<ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa>);
- vi) *Staphylococcus pyogenes* (ftp://ftp.tigr.org/pub/data/s_pneumoniae/gsp.contigs.112197.Z);
- 30 vii) PRODOM (ftp://ftp.toulouse.inra.fr/pub/prodom/current_release/prodom99_1.forblast.gz);
- viii) DOMO (<ftp://ftp.infobiogen.fr/pub/db/domo/>);

ix) TREMBL (ftp://www.expasy.ch/databases/sp_tr_nrdb/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage 44AHJD are shown in Tables 19 & 20.

5 **Example 10. Sub-Cloning of Bacteriophage 44 AHJD ORFs.**

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage 44 AHJD ORF sequence is inducible. For example, the shuttle vector pT0021, in which the firefly luciferase (*lucFF*) expression is controlled by the *ars* (arsenite) promoter/operator (Tauriainen et al., 1997), can be modified in the following fashion. Two oligonucleotides corresponding to a short antigenic peptide derived from the hemagglutinin protein of influenza virus (HA epitope tag) were synthesized (Field et al., 1988). The sense strand HA tag sequence (with *Bam*HI, *Sal*I and *Hind*III cloning sites) is:

5'-gatcccggtcgaccaagcttTACCCATACGACGTCCCAGACTACGCCAGCTGA-3'

15 (where upper case letters denote the nucleotide sequence of the HA tag); the antisense strand HA tag sequence (with a *Hind*III cloning site) is:

5'-agctTCAGCTGGCGTAGTCTGGGACGTCGTATGGGTAaagcttggtcgaccgg-3'

(where upper case letters denote the sequence of the HA tag). The two HA tag oligonucleotides were annealed and ligated into pT0021 vector which had been digested with *Bam*HI and *Hind*III. This manipulation resulted in replacement of the *lucFF* gene by the HA tag. This modified shuttle vector containing the *arsenite* inducible promoter, the *arsR* gene, and HA tag was named pTHA. A diagram outlining our modification of pT0021 to generate pTHA is shown in Fig. 1A (another useful vector construct is shown in Fig. 1B).

25 Each ORF, encoded by Bacteriophage 44 AHJD, larger than 33 amino acids and having a Shine-Dalgarno sequence upstream of the initiation codon can be selected for functional analysis for bacterial inhibition. Each individual ORF, from initiation codon to last codon (excluding the stop codon), can be amplified from phage genomic DNA using the polymerase chain reaction (PCR). For PCR amplification of
30 ORFs, each sense strand primer targets the initiation codon and is preceded by a *Bam*HI restriction site (5'-cgggatcc-3') and each antisense oligonucleotide targets the pentultimate codon (the one before the stop codon) of the ORF and is preceded by a *Sal*I restriction site (5'-gcgtcgaccg-3'). The PCR product of each ORF can be gel

purified and digested with *Bam*HI and *Sal*I. The digested PCR product can then be gel purified using the Qiagen kit as described, ligated into *Bam*HI and *Sal*I digested pTHA vector, and used to transform *E. coli* bacterial strain DH10 β (as described above). As a result of this manipulation, the HA tag is set in frame with the ORF and is positioned at the carboxy terminus of each ORF (pTHA/ORF clones). Recombinant pTHA/ORF clones will be picked and their insert sizes were confirmed by PCR analysis using primers flanking the cloning site. The following primers can be used for PCR amplification: HAF: 5'TATTATCCAAACTTGAACA3'; HAR: 5'CGGTGGTATATCCAGTGATT'. The sequence integrity of cloned ORFs can be verified directly by DNA sequencing using primers HAF and HAR. In cases where verification of ORF sequence can not be achieved by one pass with the sequencing primers, additional internal primers will be selected and used for sequencing.

Staphylococcus aureus strain RN4220 (Kreiwirth et al., 1983) will be used as a recipient for the expression of recombinant plasmids. Electoporation will be performed essentially as previously described (Schenk and Laddaga, 1992). Selection of recombinant clones will be performed on Luria-Broth agar (LB-agar) plates containing 30 μ g/ml of kanamycin.

Alternatively, a constitutive promoter can be used to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids will be introduced into *Staphylococcus aureus* strain RN4220 (Kreiwirth et al., 1983) using electoporation as previously described (Schenk and Laddaga, 1992).

Cloning of ORFs with a Shine-Dalgarno sequence

ORFs with a Shine-Dalgarno sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), can be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10. Recombinant clones are then picked and their insert sizes confirmed by

PCR analysis using primers flanking the cloning site as well as restriction digestion. The sequence fidelity of cloned ORFs can be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site internal
5 primers can be selected and used for sequencing. Recombinant plasmids can be introduced into *Staphylococcus aureus* strain RN4220 (Kreiwirth et al., 1983) using electroporation as previously described (Schenk and Laddaga, 1992).

Induction of gene expression from the *ars* promoter.

If an inducible promoter is used, e.g., the *ars* promoter, induction can be
10 assessed, for example, in either of the two methods.

1. Screening on agar plates

The functional identification of killer ORFs can be performed by spreading an aliquot of *S. aureus* transformed cells containing phage 44 AHJD ORFs onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5 μ M). The
15 plates are incubated overnight at 37°C, after which a growth inhibition of the ORF transformants on plates that contain arsenite are compared to plates without arsenite.

2. Quantification of growth inhibition in liquid medium

Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are
20 then diluted to the mid log phase ($OD_{540}=0.2$) with fresh media containing antibiotic and transferred to 96-well microtitration plates (100 μ l/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 μ M) and the culture incubated for an additional 2 hrs at 37°C. The effect of expression of the phage 44 AHJD ORFs on bacterial cell growth is then monitored by measuring the OD_{540} and comparing the
25 rate of growth to the culture not containing inducer. [As positive controls for growth inhibition, the *kilA* gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 *Virology* #193: 1033-1036), and the *holin/lys* genes of the *Staphylococcus aureus* phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G., Maier, SK. and Scherer, S. 1998. *FEMS Microbiology Letters* #162:265-274) can be
30 subcloned into the *ars* inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic-selection but lacking inducer. Following incubation overnight at 37°C, the number of

- colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing full bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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Example 11. Growth of *Enterococcus* bacteriophage 182 and purification of genomic DNA.

The *Enterococcus* propagating strain (PS) (*Enterococcus* sp. Group D, Felix d'Herelle Reference Centre #HER 1080) was used as host to propagate its respective
10 phage 182 (Felix d'Herelle Reference Centre #HER 80). Two rounds of plaque purification of phage 182 were performed on soft agar essentially as described in Sambrook *et al.* (1989). Briefly, the *Enterococcus* sp. PS strain was grown overnight at 37°C in Tryptic Soy Broth [TSB: 17 g Bacto tryptone, 3 g Bacto soytone, 2.5 g Bacto dextrose, 5 g Sodium chloride, and 2.5 g Dipotassium phosphate per liter
15 (Difco Laboratories (#0370-17-3)]. The culture was then diluted 20 fold in TSB and incubated at 37°C until the OD₅₄₀ = 0.2 (early log phase) with constant agitation. In order to obtain single plaques, phage 182 was subjected to 10 fold serial dilutions using the phage buffer (1 mM MgSO₄, 5 mM MgCl₂, 80 mM NaCl and 0.1% Gelatin (w/v)) and 10 l of each dilution was used to infect 0.5 ml of the bacterial cell
20 suspension. After incubation at 15 min at 37°C, 2 ml of melted soft agar (TSB supplemented with 0.6% agar) was added to the mixture and poured onto the surface of 100 mm Tryptic Soy Agar plates [TSA: 15 g Tryptone peptone, 5 g Soytone peptone, 5 g Sodium chloride and 15 g of Agar per liter (Difco Laboratories #0369-17)]. After overnight incubation at 37°C, a single plaque was isolated, resuspended in
25 1 ml of phage buffer by end over end rotation for 2 hrs at room temperature, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock for all subsequent manipulations.

The propagation procedure for bacteriophage 182 was modified from the agar
30 layer method of Swanstörn and Adams (1951). Briefly, the *Enterococcus* sp. PS strain was grown to stationary phase overnight at 37°C in TSB. The culture was then diluted 20 fold in TSB and incubated at 37°C until the A₅₄₀ = 0.2. The suspension (15x10⁷ Bacteria) was then mixed with 15x10⁵ plaque forming units (pfu) to give a

ratio of 100-bacteria/pfu. After incubation of 15 min at 37°C, 7.5 ml of melted soft agar (TSB plus 0.6% agar) were added to the mixture and poured onto the surface of 150 mm TSA plates and incubated 16 hrs at 37°C. To collect the plate lysate, 20 ml of TSB were added to each plate and the soft agar layer was collected by scrapping off
5 with a clean microscope slide followed by vigorous shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant fluid (lysate) is collected and subjected to a treatment with 10 µg /ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to
10 10% (w/v) of PEG 8000 and 0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO₄, 5 mM MgCl₂, 80 mM NaCl and 0.1% Gelatin). The phage suspension was extracted with 1 volume of chloroform and further purified by
15 centrifugation on a cesium chloride step gradient as described in Sambrook *et al.* (1989), using a TLS 55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 hrs at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 xg) for 24 hrs at 4°C using a TLV rotor (Beckman). The phages
20 were harvested and dialyzed for 4 hrs at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl₂. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 g/ml Proteinase K and 0.5% SDS and incubating for 1 hr at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of
25 chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

Example 12. DNA sequencing of the Bacteriophage 182 genome.

Four micrograms of phage DNA was diluted in 200 µl of TE (10 mM Tris,
30 [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4

cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0]) as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 µl of 1 mM Tris [pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of *E. coli* DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume, 100 µl) containing sonicated phage DNA, 10 mM Tris-HCl [pH 8.0], 50 mM NaCl, 10 mM MgCl₂, 1 mM DTT, 50 µg/ml BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of the Klenow large fragment of DNA polymerase I (New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was precipitated with ethanol and the final DNA pellet resuspended in 20 µl of H₂O.

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of the pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 µl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 µl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10β according to standard procedures (Sambrook *et al.*, 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 µl LB and 100 µg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *Hinc* II cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 µl reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl₂, 0.02% gelatin, 1 µM primer, 187.5 µM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec

denaturation at 94°C, 30 sec annealing at 58°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep™ spin miniprep kit (Qiagen).

5 The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye™ primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism Big Dye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and
10 the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism BigDye™ terminator cycle sequencing ready reaction kit.

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Example 13. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher™ 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI
20 prism BigDye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Enterococcus* bacteriophage 182 is shown in Table 21.

A software program was used on the assembled sequence of bacteriophage 182 to identify all putative ORFs larger than 33 codons. The software scans the primary
25 nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI([http://www.ncbi.nlm.nih.gov/htbin-](http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c)
30 [post/Taxonomy/wprintgc?mode=c](http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c)) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the

next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence. The predicted ORFs for bacteriophage 182 are listed in Tables 22 & 23. Sequence homology searches for each ORF were carried out using an implementation of BLAST programs. Downloaded public databases used for sequence analysis include:

- 10 (i) non-redundant GenBank (<ftp://ncbi.nlm.nih.gov/blast/db/nr.Z>),
- ii) Swissprot (<ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z>);
- iii) vector (<ftp://ncbi.nlm.nih.gov/blast/db/vector.Z>);
- iv) pdbaa databases (<ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z>);
- v) staphylococcus aureus NCTC 8325 ([ftp://ftp.genome.ou.edu/pub/staph/staph-](ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa)
- 15 [1k.fa](ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa));
- vi) streptococcus pyrogenes
(ftp://ftp.tigr.org/pub/data/s_pneumoniae/gsp.contigs.112197.Z);
- vii) PRODOM
(ftp://ftp.toulouse.inra.fr/pub/prodom/current_release/prodom99.1.forblast.gz);
- 20 viii) DOMO (<ftp://ftp.infobiogen.fr/pub/db/domo/>);
- ix) TREMBL (ftp://www.expasy.ch/databases/sp_tr_nrd/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage 182 are shown in Tables 24 & 26.

25 **Example 14. Sub-Cloning of Bacteriophage 182 ORFs.**

Preparation of the shuttle expression vector

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage 182 ORF sequence is inducible. For example, the plasmid pND50 replicates in *E. coli*, *E. faecalis*, and *S. aureus*

30 (Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S., and Inoue, M. 1996. *Antimicrob. Agents Chemother.* 40, 1157-1163). This plasmid can be modified by conventional techniques to insert the inducible arsenite promoter, derived from the shuttle vector pT0021, in which the firefly luciferase (*lucFF*)

expression is controlled by the *ars* promoter/operator from a *S. aureus* plasmid (Tauriainen, S., Karp, M., Chang, W and Virta, M. (1997). Recombinant luminescent bacteria for measuring bioavailable arsenite and antimonite. *Appl. Environ. Microbiol.* 63:4456-4461). This modified shuttle vector will contain the *ars* promoter, *arsR* gene
5 and a cloning site for introduction of individual phage ORFs downstream from a shine-dalgarno sequence.

Other inducible regulatory sequences can be utilized instead of the arsenite-inducible system. An example is a nisin-inducible system The *nisA* promoter activity is dependent on the proteins NisR and NisK, which constitute a two-component signal
10 transduction system that responds to the extracellular inducer nisin. The nisin sensitivity and inducer concentration required for maximal induction varies among the strains, but is functional in *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus pneumoniae*, *Enterococcus faecalis*, and *Bacillus subtilis*. Significant induction of the *nisA* promoter (10- to 60-fold induction) can be obtained in all of the
15 species. A vector containing this promoter was published as Eichenbaum Z, Federle MJ, Marra D, de Vos WM, Kuipers OP, Kleerebezem M, and Scott JR (1998) *Appl Environ Microbiol* 64, 2763-2769. Other vectors, e.g., plasmids, can also be utilized which will allow replication and transcription in *Enterococcus*.

Alternatively, a constitutive promoter can be used (e.g., the β -lactamase
20 promoter is constitutive in *E. faecalis* – see ref. 1) to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids are introduced into *E. faecalis* strain FA2-2 by electroporation, as previously described (Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S.,
25 and Inoue, M. 1996. *Antimicrob. Agents Chemother.* 40, 1157-1163).

Cloning of ORFs with a Shine-Dalgarno sequence

ORFs with a Shine-Dalgarno sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), will be amplified by PCR from phage genomic DNA. For PCR amplification
30 of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on

the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10 β . Recombinant clones are then picked and their insert sizes confirmed by PCR analysis using primers flanking the cloning site as well as restriction digestion.

- 5 The sequence fidelity of cloned ORFs will be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site internal primers will be selected and used for sequencing. Recombinant plasmids will be introduced into *E. faecalis* strain FA2-2 by electroporation, as previously described
- 10 (Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S., and Inoue, M. 1996. Antimicrob. Agents Chemother. 40, 1157-1163).

Induction of gene expression from the *ars* promoter.

If an inducible promoter is used, e.g., the *ars* promoter, induction can be assessed, for example, in either of the two methods.

15 **1. Screening on agar plates**

The functional identification of killer ORFs can be performed by spreading an aliquot of *E. faecalis* transformed cells containing phage 182 ORF onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5 μ M). The plates are incubated overnight at 37°C, after which a growth inhibition of the ORF

- 20 transformants on plates that contain arsenite are compared to plates without arsenite.

2. Quantification of growth inhibition in liquid medium

- Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are then diluted to the mid log phase ($OD_{540}=2$) with fresh media containing antibiotic
- 25 and transferred to 96-well microtitration plates (100 μ l/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 μ M) and the culture incubated for an additional 2 h at 37°C. The effect of expression of the phage 182 ORFs on bacterial cell growth is then monitored by measuring the OD_{540} and comparing the rate of growth to the culture not containing inducer. As positive controls for growth
- 30 inhibition, the *kilA* gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 *Virology* #193: 1033-1036), and the *holin/lys* genes of the *Staphylococcus aureus* phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G.,

Maier, SK. and Scherer, S. 1998. *FEMS Microbiology Letters* #162:265-274) were subcloned into the *ars* inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic selection but lacking inducer. Following incubation overnight at 37°C, the number of colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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Example 15. Growth of *Streptococcus* bacteriophage Dp-1 and purification of genomic DNA.

The *Streptococcus pneumoniae* R6 propagating strain (PS) (Tomasz, 1966) was used as host to propagate its respective phage Dp-1 (McDonnell et al., 1975). (Alternatively, *Streptococcus (Diplococcus) pneumoniae* R36A could be used. Strain R36A is available from ATCC as #11733 or 27336. *Streptococcus pneumoniae* is also available from Felix d'Herelle Reference Center in Quebec, Canada as catalog number HER 1054. Other *S. pneumoniae* strains are also available from ATCC.) Two rounds of plaque purification of phage Dp-1 were performed on soft agar essentially as described in Sambrook et al. (1989). Briefly, the *Streptococcus* R6 PS strain was grown overnight at 37°C in K-Cat media [K-Cat: 10 g Bacto casitone, 5 g Bacto tryptone, 1 g Yeast extract, 5g Potassium chloride, 0.2% Glucose, 30mM Potassium phosphate buffer [pH 8] and 250,000 Units Catalase per liter (Boehringer Mannheim #10683600). The culture was then diluted 20 fold in K-CAT and

incubated at 37°C until the $OD_{540} = 0.2$ (early log phase) with constant agitation. In order to obtain single plaques, Dp-1 phage was subjected to 10-fold serial dilutions using the phage buffer (100 mM Tris-HCl [pH 7.5], 100 mM NaCl and 10 mM $MgCl_2$) and 10 μ l of each dilution was used to infect 0.5 ml of the cell suspension.

- 5 After incubation of 15 min at 37°C, 2 ml of melted soft agar (K-CAT supplemented with 0.8% of agar) were added to the mixture and poured onto the surface of 100 mm K-CAT agar plates [K-CAT supplemented with 1.2 % of agar]. After solidification of the soft agar layer, an additional 5 ml of melted soft agar was added to visualize distinct plaques (Ronda et al., 1978). After overnight incubation at 37°C, a single
10 plaque was isolated, resuspended in 1 ml of phage buffer by end over end rotation for 2 hrs at room temperature, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock for all subsequent manipulations.

- The propagation procedure for bacteriophage Dp-1 was modified from the
15 agar layer method of Swanstörn and Adams (1951). Briefly, the R6 strain of *Streptococcus pneumoniae* was grown to stationary phase overnight at 37°C in K-CAT. The culture was then diluted 20 fold in K-CAT and incubated at 37°C until the $OD_{540} = 0.2$. The suspension (15×10^7 Bacteria) was then mixed with 15×10^5 plaque forming units (pfu) to give a ratio of 100-bacteria/pfu. After incubation of 15 min at
20 37°C, 7.5 ml of melted soft agar (K-CAT plus 0.8% agar) were added to the mixture and poured onto the surface of 150 mm K-CAT agar plates and incubated 16 hrs at 37°C. After solidification of the soft agar layer, 7.5 ml of melted soft agar were added to each plate. To collect the plate lysate, 20 ml of K-CAT media were added to each plate and the soft agar layers were collected by scrapping off with a clean microscope
25 slide followed by vigorous shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant (lysate) was collected and subjected to a treatment with 10 μ g /ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) of PEG 8000 and
30 0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (100 mM Tris-HCl [pH 7.5], 100 mM NaCl and 10 mM $MgCl_2$). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a
35 cesium chloride step gradient as described in Sambrook et al. (1989), using a TLS-55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 hrs at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an

isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 xg) for 24 hrs at 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 hrs at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl₂. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 µg/ml Proteinase K and 0.5% SDS and incubating for 1 hr at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1 mM EDTA).

Example 16. DNA sequencing of the Bacteriophage Dp-1 genome.

Four micrograms of phage DNA was diluted in 200 µl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 sec spaced by 15 sec cooling in ice/water for 3 to 4 cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0]) as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 µl of 1 mM Tris [pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of *E. coli* DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume, 100 µl) containing sonicated phage DNA, 10 mM Tris-HCl [pH 8.0], 50 mM NaCl, 10 mM MgCl₂, 1 mM DTT, 50 µg/ml BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of the Klenow large fragment of DNA polymerase I (New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was precipitated with ethanol and the final DNA pellet resuspended in 20 µl of H₂O.

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of the pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 µl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 µl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection

of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10 β according to standard procedures (Sambrook *et al.*, 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 μ l LB and 100 μ g/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *Hinc* II cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 μ l reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl₂, 0.02% gelatin, 1 μ M primer, 187.5 μ M each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 58°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep™ spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye™ primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism Big Dye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism Big Dye™ terminator cycle sequencing ready reaction kit.

Example 17. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher™ 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI prism BigDye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Streptococcus* bacteriophage Dp-1 is shown in Table 28.

A software program was used on the assembled sequence of bacteriophage Dp-1 to identify all putative ORFs larger than 33 codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG,

GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI(<http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c>) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence. The predicted ORFs for bacteriophage Dp-1 are listed in Tables 29 and 30, and Fig. 6.

Sequence homology searches for each ORF were carried out using an implementation of BLAST programs. Downloaded public databases used for sequence analysis include:

- (i) non-redundant GenBank (<ftp://ncbi.nlm.nih.gov/blast/db/nr.Z>),
- ii) Swissprot (<ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z>);
- iii) vector (<ftp://ncbi.nlm.nih.gov/blast/db/vector.Z>);
- iv) pdbaa databases (<ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z>);
- v) staphylococcus aureus NCTC 8325
(<ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa>);
- vi) streptococcus pyogenes
(ftp://ftp.tigr.org/pub/data/s_pneumoniae/gsp.contigs.112197.Z);
- vii) PRODOM
(ftp://ftp.toulouse.inra.fr/pub/prodom/current_release/prodom99.1.forblast.gz);
- viii) DOMO (<ftp://ftp.infobiogen.fr/pub/db/domo/>);
- ix) TREMBL (ftp://www.expasy.ch/databases/sp_tr_nrd/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage Dp-1 are shown in Table 31.

Example 18. Sub-Cloning of Bacteriophage Dp-1 ORFs.

Preparation of the shuttle expression vector

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage Dp-1 ORF sequence is inducible. For example, the plasmid pLSE4 replicates in *E. coli*, and *S. pneumoniae* (Diaz and Garcia, 1990). This plasmid can be modified by conventional techniques to insert the inducible arsenite promoter, derived from the shuttle vector pT0021, in which the

firefly luciferase (*lucFF*) expression is controlled by the *ars* promoter/operator from a *S. aureus* plasmid (Tauriainen, S., Karp, M., Chang, W and Virta, M. (1997).

Recombinant luminescent bacteria for measuring bioavailable arsenite and antimonite. *Appl. Environ. Microbiol.* 63:4456-4461). This modified shuttle vector will contain the *ars* promoter, *arsR* gene and a cloning site for introduction of individual phage ORFs downstream from a shine-dalgarno sequence.

Other inducible regulatory sequences can be utilized instead of the arsenite-inducible system. An example is a nisin-inducible system. The *nisA* promoter activity is dependent on the proteins NisR and NisK, which constitute a two-component signal transduction system that responds to the extracellular inducer nisin. The nisin sensitivity and inducer concentration required for maximal induction varies among the strains, but is functional in *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus pneumoniae*, *Enterococcus faecalis*, and *Bacillus subtilis*. Significant induction of the *nisA* promoter (10- to 60-fold induction) can be obtained in all of the species. A vector containing this promoter was published as Eichenbaum Z, Federle MJ, Marra D, de Vos WM, Kuipers OP, Kleerebezem M, and Scott JR (1998) *Appl Environ Microbiol* 64, 2763-2769. Other vectors, e.g., plasmids, can also be utilized which will allow replication and transcription in *Streptococcus*.

Alternatively, a constitutive promoter can be used to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids are introduced into *S. pneumoniae* R6 as previously described (Diaz and Garcia, 1990)

Cloning of ORFs with a Shine-Dalgarno sequence

ORFs with a Shine-Dalgarno sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), will be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10 β . Recombinant clones are then picked and their insert sizes confirmed by PCR analysis using primers flanking the cloning site as well as restriction digestion. The sequence fidelity of cloned ORFs will be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site

internal primers will be selected and used for sequencing. Recombinant plasmids will be introduced into *S. pneumoniae* R6 as previously described (Diaz and Garcia, 1990).

Induction of gene expression from the *ars* promoter.

- 5 If an inducible promoter is used, e.g., the *ars* promoter, induction can be assessed, for example, in either of the two methods.

1. Screening on agar plates

- The functional identification of killer ORFs can be performed by spreading an aliquot of *S. pneumoniae* transformed cells containing phage Dp-1 ORFs onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5 μ M).
10 The plates are incubated overnight at 37°C, after which a growth inhibition of the ORF transformants on plates that contain arsenite are compared to plates without arsenite.

2. Quantification of growth inhibition in liquid medium

- Cells containing different recombinant plasmids can be grown for overnight at
15 37°C in LB medium supplemented with the appropriate antibiotic selection. These are then diluted to the mid log phase ($OD_{540}=2$) with fresh media containing antibiotic and transferred to 96-well microtitration plates (100 μ l/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 μ M) and the culture incubated for an additional 2 hrs at 37°C. The effect of expression of the phage Dp-1 ORFs on
20 bacterial cell growth is then monitored by measuring the OD_{540} and comparing the rate of growth to the culture not containing inducer. [As positive controls for growth inhibition, the *kilA* gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 *Virology* #193: 1033-1036), and the *holin/lysin* genes of the
Staphylococcus aureus phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G.,
25 Maier, SK. and Scherer, S. 1998. *FEMS Microbiology Letters* #162:265-274) can be subcloned into the *ars* inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic selection but lacking inducer. Following incubation overnight at 37°C, the number of colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but
30 detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing full bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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10 All patents and publications mentioned in the specification are indicative of the levels of skill of those skilled in the art to which the invention pertains. All references cited in this disclosure are incorporated by reference to the same extent as if each reference had been incorporated by reference in its entirety individually.

15 One skilled in the art would readily appreciate that the present invention is well adapted to carry out the objects and obtain the ends and advantages mentioned, as well as those inherent therein. The specific methods and compositions described herein as presently representative of preferred embodiments are exemplary and are not intended as limitations on the scope of the invention. Changes therein and other uses will occur to those skilled in the art which are encompassed within the spirit of the invention are defined by the scope of the claims.

20 It will be readily apparent to one skilled in the art that varying substitutions and modifications may be made to the invention disclosed herein without departing from the scope and spirit of the invention. For example, those skilled in the art will recognize that the invention may suitably be practiced using a variety of different bacteria, bacteriophage, and sequencing methods within the general descriptions provided.

25 The invention illustratively described herein suitably may be practiced in the absence of any element or elements, limitation or limitations which is not specifically disclosed herein. Thus, for example, in each instance herein any of the terms "comprising," "consisting essentially of" and "consisting of" may be replaced with either of the other two terms. The terms and expressions which have been employed 30 are used as terms of description and not of limitation, and there is not intention that in the use of such terms and expressions of excluding any equivalents of the features shown and described or portions thereof, but it is recognized that various modifications are possible within the scope of the invention claimed. Thus, it should 35 be understood that although the present invention has been specifically disclosed by preferred embodiments and optional features, modification and variation of the concepts herein disclosed may be resorted to by those skilled in the art, and that such modifications and variations are considered to be within the scope of this invention as defined by the appended claims.

In addition, where features or aspects of the invention are described in terms of Markush groups or other grouping of alternatives, those skilled in the art will recognize that the invention is also thereby described in terms of any individual member or subgroup of members of the Markush group or other group. For example, 5 if there are alternatives A, B, and C, all of the following possibilities are included: A separately, B separately, C separately, A and B, A and C, B and C, and A and B and C. Thus, for example, for the bacteria and phage specified herein, the embodiments expressly include any subset or subgroup of those bacteria and/or phage. While each such subset or subgroup could be listed separately, for the sake of brevity, such a 10 listing is replaced by the present description.

Thus, additional embodiments are within the scope of the invention and within the following claims.

Table 1**Phages against human and animal pathogenic bacteria**

5

I. Pathogen name	Phage name	II. Catalog#	Origin/reference
<i>Acinetobacter calcoaceticus</i>	A3/2 A10/45 A36 B9GP B ₉ PP BS46 E13 E14 531		Felix d'Herelle Reference Centre, Quebec, Quebec
	Ap3 P78		J. Bacteriol 1984. 157: 179-183 J. Gen. Microbiol 1986.132: 2633-2636
<i>Acinetobacter haemolyticus</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Acinetobacter johnsonii</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Acinetobacter sp.</i>	BP1		J. Virol. 1968.2:716-722
	G4, HP2, HP3 & HP4		Can. J. Microbiol. 1966.12:1023-1030 & J. Virol. 1974.13:46-52 & Arch. Virol. 1994.135:345-354
	A1, A4, A9 & 196		Arch. Virol. 1994.135:345-354
	HP1		Can. J. Microbiol. 1966.12:1023-1030
	A19, A23, A29, A31, A33, A34, A3759 & 2845		J. Microsc (Paris) 1973.16:215-224 & CR. Hebdo Seances Acad. Sci. Ser D. Sci Natur (Paris) 278:1907-1909 & Arch. Virol. 1994.135:345-354 & Rev. Can. Biol. 1970.29:317-320
<i>Actinobacillus actinomycetecomitans</i>			FEMS Microbiol Lett 1994. 119:329-337

			Infec. Immun. 1982. 35: 343-349
			Mol.Gen.Genet 1998.258: 323-325
	Aap247		Oral Micriol. Immunol 1997.12: 40-46
<i>Actinomyces viscosus</i>	43146-B1		The American Type Culture Collection
			Infect.Immun.1985.48:228-233
			Infect.Immun.1988.56:54-59
			Plasmid 1997.37:141-153
<i>Aeromonas hydrophila</i>	PM2** & PM3		FEMS Microbiol.Lett. 1990.57:277-282
	Aeh1 Aeh2 PM4 PM5 PM6 T7-ah		Felix d'Herelle Reference Centre,Quebec,Quebec

<i>Aeromonas salmonicida</i>	3 25 29 31 32 40RR _{2.8} t 43 51 56 59.1 65 Asp37		Felix d'Herelle Reference Centre, Quebec, Quebec
	55R.1		Can. J. Microbiol. 1983. 29: 1458-1461
<i>Alteromonas espejiana</i>	PM2**	27025-B1	The American Type Culture Collection
<i>Asticcacaulis biprosthecum</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Asticcacaulis excentricus</i>		15261-B1 15261-B2 15261-B3	The American Type Culture Collection
	φAc21 φAc24		
<i>Azotobacter vinelandii</i>	A14 A21 A31 A41 PAV1	12518-B1 12518-B4 12518-B5 12518-B9 12518-B10 13705-B1	The American Type Culture Collection
<i>Azotobacter sp.</i>			Virology 1972.49:439-452
<i>Bacteroides fragilis</i>	Bf-1		Rev. Infect. Dis. 1979. 1: 325-336
	B40-8		FEMS Microbiol. Lett. 1991. 66: 61-67
	HSP40		Appl. Environ. Microbiol. 1989. 55: 2696-2701
	phiA1		Zentralbl. bakteriol. 1972. 222: 57-63
<i>Bdellovibrio bacteriovorus</i>	MAC-1		J. Gen. Microbiol. 1987. 133: 3065-3070
<i>Bdellovibrio sp.</i>	VL-1		J. Virol. 1973. 12: 1522-1533
<i>Bordetella brochiseptica</i>	214		Zh. Mikrobiol. Epidemiol. Immuno. 1987. 5: 9-13

<i>Bordetella parapertussis</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
			Mol. Gen. Mikrobiol. Virusol. 1988.4: 22-25
			Zh.Mikrobiol.Epidemiol.Immuno. 1987.5:9-13
	41405		Zh.Mikrobiol.Epidemiol.Immuno. 1987.5:9-13
<i>Brucella abortus</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
	10/I 24/II 212/XV	23448-B1 23448-B2 23448-B3 17385-B1 17385-B2	The American Type Culture Collection
	BK-2, TB & Fi**		Zh.Mikrobiol.Epidemiol.Immunobiol.1983.2: 48-52
	R/c & R/O		Dev. Biol. Stand. 1984.56: 55-62
	R/c		Dev. Biol. Stand. 1984.56: 55-62
<i>Brucella canis</i>	BK-2	23456-B1	The American Type Culture Collection
<i>Brucella melitensis</i>	Wb		Zentralbl. Veterinarmed.1975.22:866-867
<i>Brucella suis</i>			

	Fi** & TB		Zh.Mikrobiol.Epidemiol.Immunobiol.1983.2: 48-52
<i>Brucella sp.</i>			Can. J. Vet. Res. 1989.53: 319-325
			Res. Vet. Sci. 1988. 44: 45-49
	R		Zh.Mikrtobiol.Epidemiol.Immunobiol.1983.2: 48
<i>Campylobacter coli</i>		43133-B1	The American Type Culture Collection
<i>Campylobacter coli</i> (Cont'd)	18	43134-B1	The American Type Culture Collection
	19	43135-B1	
	20	43136-B1	
<i>Campylobacter jejuni</i>	1	35918-B1	The American Type Culture Collection
	2	35919-B1	
	3	35920-B1	
	4	35921-B1	
	5	35918-B2	
	6	35920-B2	
	7	35922-B2	
	8	35923-B1	
	9	35924-B1	
	10	35925-B1	
	11	35925-B2	
	12	35922-B2	
	13	35924-B2	
	14	35922-B3	
	17	43133-B1	
	18	43134-B1	
	19	43135-B1	
	20	43136-B1	
<i>Campylobacter</i> (<i>Helicobacter</i>) <i>pylori</i>	HP1		J. Med. Microbiol.1993. 38: 245-249
<i>Chlamydia psittaci</i>	Chp1**		J. Gen. Virol. 1989. 70: 3381-3390
<i>Clostridium acetobutylicum</i>	CAK-1		J.Bacteriol.1993.175:3838-3843

<i>Clostridium botulinum</i>			Nucleic Acids Res.1990.18:1291
			Bioch.Biophys.res.Commun.1990.171.1304-1311
			Microbiol.immunol.1981.25:915-927
			J.Vet.Med.Sci.1992.54:675-684
	CE β & CE γ		
<i>Clostridium difficile</i>	41 & 56		J. Clini.Microbiol. 1985.21:251-254

<i>Clostridium perfringens</i>			Rev.Can.Biol.1977.36:205-215
			FEMS Microbiol.Lett. 1990.54:323-326
<i>Clostridium sporogenes</i>	59 70 71 72S 72L	8074-B1 17886-B1 17886-B3 17886-B4 17886-B5 17886-B6	The American Type Culture Collection
<i>Clostridium tetani</i>	A & B		Rev.Can.Biol.1978.37:43-46
<i>Corynebacterium diptheriae</i>			Vopr.Virusol.1986.31:577-584
<i>Corynebacterium pseudotuberculosis</i>	NN	12319-B1	The American Type Culture Collection
<i>Corynebacterium sp</i>	DLC 2921/49	12052-B1	The American Type Culture Collection

<i>Enterococcus faecalis</i>	42	19948-B1	The American Type Culture Collection
<i>Enterococcus faecium</i>	124 133	19950-B1 19953-b2 19953-B1	The American Type Culture Collection

<i>Escherichia coli</i>		11303-B14 11303-B10 11303-B21 8677-B1 11303-B13 13706-B4	The American Type Culture Collection
<i>Escherichia coli</i> (Cont'd)		15766-B1 15766-B1 1242-B5 15669-B2 15767-B1 11303-B16 27-65-B1 25065-B2 15669-B1 15597-B1 21816-B1 23724-B9 15593-B1 25404-B1 29746-B1 23631-B1 25868-B1 25298-B1 25298-B2 11303-B37 11303-B24 11303-B26 11303-B27 11303-B28 11303-B29 11303-B30 11303-B33 11303-B31 11303-B25 11303-B35 11303-B34 11303-B36 11303-B32 13706-B5 11303-B1 11303-B2 11303-B3 11303-B4 35060-B1 35060-B2 35060-B3 11303-B5 11303-B6 11303-B7 11303-B38 12141-B1	The American Type Culture Collection
	C204 E1 f1** f2** FCZ fd**		
	If1**		
	MS2** MU9 Mu-1 Ox6 P1** P4 sid _i ** Q-β** R17** Z1K/1 ZJ/2		

<i>Escherichia coli</i> (Cont'd)		11303-B20	The American Type Culture Collection
		11303-B17	
		11303-B15	
		11303-B11	
	547	11303-B18	
	UV1	13706-B2	
	UV47	23724-B2	
	UV375	23724-B1	
	$\alpha 3^{**}$	23724-B3	
	λ^{**}	23724-B4	
	λ C-17	23724-B5	
	λ sus P-3	23724-B6	
	λ sus R-5	23724-B7	
	λ sus J-6	23724-B8	
	λ sus O-8	35860-B1	
	λ sus A-11	13706-B3	
	λ ind'	15597-B2	
	$\phi 92$	13706-B1	
	ϕR	49696-B1	
	$\phi V-1$		
	$\phi X174^{**}$		
	$\phi Xcs70am-3$		
	G4** & ϕK^{**}		Biochim.Biophysica Acta.1992.1130:277-288
	BF23**		J.Bacteriol.1977.129:265-275
	Mu1		J.Ultrastruct.Res.1966.14:441-448
	Hp17		J.Mol.Biol.1991.218:705-721
	K3** & Ox2**		FEBS Lett.1987.215:145-150
	Rb18**, Rb51 & Rb69**		J.Bacteriol.1990.172:180-186
	H1**, H3, H8, K9, K18 & Ox1		Mol.Gen.Genet.1990.221:491-494
	M1**, Tula** & Tu1b**		J.Mol.Biol.1987.196:165-174
	K10		J.Bacteriol.1979.140:680-686
	Qsr'		J.Bacteriol.1985.162:256-262
	B278		J.Gen.Microbiol.1988.134:1333-1338
	phi 80**		FEMS Microbiol.Lett.1994.119:71-76
	phi m173		Genetika 1985.21:673-675
	tf-1		J.Gen.Microbiol.1987.133:953-960
	P4 & phiR73		Mol.Microbiol.1995.18:201-208
	I ₂ -2		J.Gen.Microbiol.1982.128:2797-2804
	PRD1		Virology 1990.177:445-451
	K3hx		Mol.Gen.Genet.1987.206:110-115
	933J** & 933W**		Infect.Immunity.1986.53:135-140
	H19-B**		J.Bacteriol.1987.169:4308-4312
	Tcp-111		Zentralbnl.Bakteriol.Mikrobiol.Hyg.1988.270:41-51

	N4**	Vet.Microbiol.1992.30:203-212
	Phi 80 trp	Ann.Inst.Pasteur.1971.120:121-125
	Obeta 1	J.Bacteriol.1978.133:172-177
	P1CM	J.Gen.Microbiol.1978.107:73-83
	PA-2**	J.Bacteriol.1990.172:1660-1662
	186**	Mol.Gen.Genet.1982.187:87-95
	186.IX.B	Mol.Microbiol.1992.6:2629-2642
	21**	Virology 1983.129:484-489
	P4**	MicrobiolRev.1993.57:683-702
	82**	J.Biol.Chem.1987.262:11721-11725
	PSP3	J.Bacteriol.1996.178:5668-5675
	HK022**	Nucleic Acids Res.1994.22:354-356
	D108**	Nucleic Acids Res.1986.14:3813-3825
<i>Escherichia coli</i> (Cont'd)	Rb49	J.Mol.Biol.1997.267:237-249
	Ike**	J.Mol.Biol.1985.181:27-39
	P22dis	Mol.Gen.Genet.1978.166:233-243
	N15**	J.Bacteriol.1996.178:1484-1486
	If1**	Proc.R.Soc.Lond.B.Biol.Sci.1991.245:23-30
	Stx2Phi-I & Stx2Phi-II	Infect.Immun.1998.66:4100-4107
	18	Virology 1987.156:122-126
	X	J.Gen.Microbiol.1981.126:389-396
	AC3	Mol.Microbiol.1991.5:715-725

	BW-1 C-1 E920g Esc-7-11 H19J Haiti HK243 Ia K20 K30 KL ₃ M Mu** O103 O157:H7 PID ptI PilH α PR64FS PR772 SS4 β 4Q λ vir** Ω 8 09-1 92		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Haemophilus influenzae</i>	HP1**		Nucleic Acids Res. 1996.24:2360-2368
	S2**		Gene 1997. 196: 139-144
<i>Halobacterium cutirubrum</i>	S45		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Halobacterium halobium</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
			Can.J.Microbiol.1982.28:916-921
<i>Halobacterium salinarum</i>			Biol.Chem.Hoppe Seyler 1994.375:747-757

<i>Klebsiella oxytoca</i>	tf-1		J.Gen.Microbiol.1987.133:953-960
<i>Klebsiella pneumoniae</i>	60	23356-B1	The American Type Culture Collection
	92	23357-B1	
	K19Q		
	FC3-1 & FC3-9		
	FC3-10		Can.J.Microbiol.1991.37:270-275
<i>Klebsiella sp.</i>	K11**		FEMS Microbiol.Lett.1991.67:291-297
<i>Leptospira sp.</i>	LE1, LE3 & LE4		Mol.Gen.Genet. 1990.221:283-286
			Res.Microbiol.1990.141:1131-1138
<i>Listeria monocytogenes</i>	243	23074-B1	The American Type Culture Collection
	197,1313 & 9425		Appl.Environ.Microbiol.1997.63:3374-3377
	H387 & H387-A		Appl.Environ.Microbiol.1993.59:2914-2917
	5775,6223 &12682		APMIS.1993.101:160-167
	2389, 2671, 4211 & 2685		Intervirology 1994.37:31-35 & Zentralbl.Bakteriol.Mikrobiol.Hyg.1986.261:1 2-28
	4b, 4ab, 4g & 3c		Ann.Microbiol (Paris) 1977.128:185-198
	A118, A500 & A511**		Mol.Microbiol. 1995.16:1231-1241-992
	1, 3, 4, 5, 6, 7, 8, 9, 10, 11, 14, 15, 16, 17, 19 & 20		Ann.Microbiol. (Paris) 1979.130B:179-189
	1/2a, 1/2b, 3c, 4ab, 6a & 6b		Clin.Invest.Med.1984.7:229-232
	φLMUP35 2685		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Listeria innocua</i>	4211		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Micrococcus luteus</i>		4698-B1	The American Type Culture Collection
		4698-B4	
	N3	4698-2	
	N4	4698-B3	
	N8		
<i>Micrococcus luteus</i>	N17		Can.J.Microbiol. 1979.25:1027-1035
<i>Mycobacterium smegmatis</i>	BK-3	27203-B1	The American Type Culture Collection
	Bo1**	27204-B1	
	Bo 6	27205-B1	
	Bo 6II	27205-B2	
	Bo 6III	27205-B3	
	Mc-2	607-B6	
	Mc-4	607-B7	
	NN	11727-B1	
	Phagus lacticola	11759-B1	
	R1	607-B1	

Legendre Leo Roy Sedge	HER 317 HER 330 HER 333 HER 335 HER 334 HER 331 HER 316	Felix d'Herelle Reference Centre, Quebec, Quebec
		Mol.Microbiol.1993.7:395-405
		J.Mol.Biol.1998.279:143-164
		Proc.Natl.Acad.Sci USA.1988.84:2833-2837
		Mol.Biol.Rep. 1981.30:11-15
		Proc.Natl.Acad.Sci.USA 1997.94:10961-10966
	29M, 31M, 122, 154, 37, 29D, 46, 139,110, 141, 74D, AG1 & DS6A	Arch.Virol.1993.133:39-49 & Am.Rev.Respir.Dis.1975.112:17-22
<i>Mycobacterium fortuitum</i>	Bo 4 Bo 7	23052-B1 27207-B1 27207-B2 The American Type Culture Collection

<i>Mycobacterium leprae</i>			Ann.Microbiol. (Paris) 1982.133:93-97
<i>Mycobacterium tuberculosis</i>	DS6A	25618-B1 25618-B2 4243-B1	The American Type Culture Collection
	110, 139 & 33D		Arch.Virol.1993.133:39-49
	AG1,GS4E, BG1, PH & BK1		The Biology of Mycobacteria.Academic Press,Toronto 1982 (Ratledge & Stanford) 1982.309-351
<i>Mycobacterium sp</i>	Phagus pellegrini NN B1	11760-B1 11761-B1 23239-B1	The American Type Collection Culture

	TM4, ph60, ph72, PhAE39, phAE40 & Bxb1		Microbiology 1995.141:1173-1181
	C2		Experientia 1969.25:1112-1113
	18 & I15		J.Gen.Virol.1987.68:949-956
	63		Gruzlica 1968.36:617-622
	phlei & butyricum		J.Gen.Virol.1975.29:235-238
	MyF3P-59a		Z.Allg.Mikrobiol.1968.8:29-37
	Bo2a		J.Gen.Virol.1973.20:75-87
	D4,D28 & D32		J.Exptl.Med.1966.123:327-340
	HC		J.Bacteriol.1963.86:608-609
<i>Mycobacterium vaccae</i>	B5	15483-B1	The American Type Culture Collection
<i>Mycobacterium phlei</i>	NN Bo 2 Bo 2h Bo 3	11728-B1 11758-B1 27086-B2 27086-B1	The American Type Culture Collection
<i>Mycoplasma arthritidis</i>	MAV1**		Infect.Immunity.1995.63:4016-4023
<i>Mycoplasma hyorhinis</i>	Hr-1		Arch.Virol.1983.77:81-85
<i>Mycoplasma pneumoniae</i>	Br-1		Arch.Virol.1983.75:1-15
<i>Mycoplasma pulmonis</i>			Plasmid 1995. 33: 41-49
<i>Mycoplasma sp.</i>			J.Gen.Microbiol.1985:131:3117-3126
			J. Virol.1986.59:584-590
			Gene 1994. 141: 1-8

		Microbios 1990. 64: 111-125
		Infection& Immunity 1995. 63: 4016-4023
		Med.Biol.1982.60:116-120
MV-L2 &		Arch.Virol.1979.61:289-296
		Acta.Virol.1978.22:443-450
		J.Gen.Virol.1979.42:315-322
		Virology 1973.55:118-126

			Science 1971.173:725-727
<i>Neisseria perflava</i>			J.Clin.Microbiol.1976. 4:87-91
<i>Nocardia erythropolis</i>	φC		J.Gen.Virol.1974.23:247-254
	φEC		J.Bacteriol.1976.126:1104-1107
<i>Pasteurella multocida</i>	B225		Arch.Exp.Veterinarmed.1981.35:433-436
	B939a		Am.J.Vet.Res.1978.39:1565-1566
	Nos.115, 32, 967 & 1075		Vet.Med.Nauki. 1977.14:33-36
<i>Propionibacterium acnes</i>	NN	29399-B1	The American Type Collection Culture

<i>Pseudomonas aeruginosa</i>	2	12175-B1	The American Type Culture Collection
	2A	12175-B2	
	2B	12175-B3	
	11	12175-B4	
	16	14205-B1	
	24	14206-B1	
	27	14207-B1	
	44	14208-B1	
	73	14209-B1	
	95	14210-B1	
	109	14211-B1	
	113	14212-B1	
	249	14213-B1	
	B3	14214-B1	
	Hoff 2	15692-B1	
	Hoff 3	14203-B1	
	Pa	14204-B1	
	Pb	12055-B1	
	PB-1	12055-B2	
	Pc	15692-B3	
	Pf	12055-B3	
	PP7**	25102-B1	
		15692-B2	
	7 & 31		Felix d'Herelle Reference Centre, Quebec, Quebec
	PF3**		J.Virol.1983.47:221-223
	φ-MC		Can.J.Microbiol.1969.15:1179-1186
	PF1**		J.Mol.Biol.1991.218:349-364
	PR4**		J.Gen.Virol.1979.43:583-592
	A7		J.Bacteriol.1992.174:2407-2411
	KF1		J.Biochem.1983.93:61-71
	φCTX**		Mol.Microbiol.1993.4:1703-1709
	φ2**		J.Virol.1977.24:135-141

	<p>φKZ, 21, φNZ, PMN17, PTB80, 68, PB-1, E79, 16, 109, 352, 1214, F8, 71, 337, M4, φC17, SL2, B17, Li-24, φmnP78, PS17**, φ1, 73, M6, Li-2, 7, φmnF82, PTB2, PTB20, PTB42, φKF77, 31, PTB21, 119x, φPLS27, B3, 258, Hw12, PM57, PM62, PM105, 148, PM681, 198, 218, 222, 242, 246, PC131, φC11, SL5, D3112**, Jb19, F7, PM69, PM13, PM61, PM113, φ240, 249 & 269</p>		ddd
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<i>Pseudomonas aeruginosa</i> (Cont'd)	297, 309, 318, 11,		Arch. Virol. 1993.131:141-151
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<i>Pseudomonas cepacia</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Pseudomonas fragi</i>	wy	27362-B1 27363 B1	The American Type Culture Collection
<i>Pseudomonas phaseolicola</i>	φ6		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Pseudomonas putida</i>	gh-1	12633-B1	The American Type Culture Collection
<i>Pseudomonas syringae</i>	φ-6	40492-B1 21781-B1	The American Type Culture Collection
<i>Pseudomonas sp.</i>	PPs-G3	49780-B1	The American Type Culture Collection
<i>Salmonella bareilly</i>	Sab 2		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Salmonella enteritidis</i>	1, 2, 3 & 6		Epidemiol. Infect. 1995.114:227-236
	2a, 3a, 4a, 5a, 6a, 7a, 8a, 9a, 15, 19, 20 & 21**		Vet. Med. Nauki. 1975.12:55-60
<i>Salmonella newington</i>	Epsilon 34		J. Struct. Biol. 1995.115:283-289
<i>Salmonella newport</i>		27869-B1 27869-B2	The American Type Culture Collection
	16-19		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Salmonella paratyphi</i>		19940-B1 12176-B1	The American Type Culture Collection
	Paratyphoid A Jersey		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Salmonella senftenberg</i>	SasL1, SaL2, SaL3, SaL4, SaL5 & SaL6		Indian J. Med. Res. 1997.105:47-52
<i>Salmonella typhimurium</i>	P22** SL-1	19585-B1 40282	The American Type Culture Collection
	MB78**		J. Virol. 1982.41: 1038-1043
	SE1		J. Gen. Microbiol. 1986.132:1035-1041
	LT2		Virology 1971.45:835-636
	ES18**		Virology 1970.42:621-632
	L**		J. Virol. 1985.56:1034-1036

	PICM clr-100		Mol.Gen.Genet.1975.138:113-126
	F22		Genet.Res.1986.48:139-143
	Fels 1		J.Gen.Virol.1978.38:263-272
	Fels 2		Genet.Res.1986.48:139-143
	Px		Mol.Gen.Genet.1970.108:184-202
	P1kc		Virology 1974.60:503-514
	A3 & A4		J.Bacteriol. 1987.169:1003-1009
	HT		Genet.Res.1976.27:315-322
<i>Salmonella typhimurium</i> (Cont'd)	IRA		J.Basic Microbiol. 1990.30:707-716
	MudI		Mol.Gen.Genet. 1986.202:327-330
	P22 (cir4-1, cir5-1 & cir6-1)		Mol.Gen.Genet.1984.198:105-109
	BF23**		Mol.Gen.Genet.1976.147:195-202
	Kb1		J.Bacteriol.1974.117:907-908
	P221dis		J.Gen.Virol.1978.41:367-376
	PRD1**		Virology 1990.177:445-451
	I _T -2**		J.Gen.Microbiol.1982.128:2797-2804
	tf-1		J.Gen.Microbiol.1987.133:953-960
<i>Salmonella typhosa/typhi</i>	X**		J.Gen.Microbiol.1981.126:389-396
	8	19937-B1	The American Type Culture Collection
	23	19938-B1	
	25	19939-B1	
	46	19942-B1	
	53	19943-B1	
	163	19946-B1	
	175	19947-B1	
	Vii	27870-B1	
	VivI	27870-B2	
<i>Salmonella sp.</i>	O1		Felix d'Herelle Refrence Centre, Quebec, Quebec
	Viii		Chung Hua Liu Hsing Ping H.T.C.1992.13:288
	j2		J.Gen.Microbiol.1983.129:3395-33400
	P3	25957-B1	The American Type Culture Collection
	P4**	25957-B2	
	P9a	25957-B3	
	P9c	25957-B4	
	P10	25957-B5	
	102	19945-B1	
	Chi (χ)	9842-B1	
<i>Sphaerotilus natans</i>	R34	97541	
	MG40		Virology 1968.34:521-530
	P14		Microb.Pathog.1990.8:393-402
	PSP3		Virology 1992.188:414
	Ike**		Zentralbl.Bakteriol.1976.234:294-304
	P27 & 9NA		J.Virol.1986.12:921-931
	SN1		Appl.Environ.Microbiol.1979.37:1025-1030

<i>Shigella dysenteriae</i>	P2 ø80	23351-B1 11456b 11456a-B1	The American Type Culture Collection
<i>Shigella flexeneri</i>	D20	12661-B1	The American Type Culture Collection
	SfII**		Mol.Microbiol.1997.26:939-950
	SfV**		Gene 1997.22:217-227
	Sf6**		Mol.Microbiol.1995.18:201-208
	SfX		Gene 1993.129:99-101
<i>Shigella sonnei</i>	C16**		
	Ufa		Mol..Biol (Mosk) 1977.11:323-331
<i>Shigella sp</i>	37	23354-B1	The American Type Culture Collection
<i>Spiroplasma citri</i>	SpV1		Plasmid 1993.29:193-205
<i>Spiroplasma sp.</i>	SpV1-R8A2B		Nucleic Acids Res. 1990.18:1293
	SpV3		Isr.J.Med.Sci.1987.23:429-433
	Sp V4		J.Bacteriol.1987.169:4950-4961
<i>Staphylococcus albus</i>			Staphylococci & Staphylococcal Infections.1997. Vol1:503-508 (Karger,Basel)

<i>Staphylococcus aureus</i>		27702-B1	The American Type Culture Collection
		27703-B1	
		27704-B1	
		23360-B1	
		23361-B1	
	15	27705-B1	
	17	27712-B1	
	29	27690-B1	
	42D**	27691-B1	
	42E	27692-B1	
	47	27693-B1	
	52	27694-B1	
	52A	27695-B1	
	53	27696-B1	
	54	27697-B1	
	55	27698-B1	
	71	27699-B1	
	75	27693-B2	
	77	27700-B1	
	79	27701-B1	
	80	27706-B1	
	81	27707-B1	
	83A	27708-B1	
	84	33742	
	85**	33741-B1	
	88	15565	
	92	19685-B1	
	5504'	11987-B1	
	K	11988-B1	
	P1	15752-B1	
	P14		
	UC18		

	HER 101 HER 239 HER 283 HER 49	Felix d'Herelle Reference Centre, Quebec, Quebec
	Twort**	
	$\phi 11^{**}$	J.Bacteriol.1988.170:2409-2411
	$\phi 13^{**}$ & $\phi 42^{**}$	J.Gen..Microbiol.1989.135:1679-1697
	L54a**	J.Bacteriol.1986.166:385-391
	80 α^{**}	Can.J.Microbiol.1996.43:612-616
	94,95 & 96	J.Clin.Microbiol.1988.26:2395-2401
	$\phi 131, A_3$ & A_5	Staphylococci & Staphylococcal Infections.1997. Vol1:503-508 (Karger,Basel)
	Phi PVL**	Gene 1998.215:57-67
<i>Staphylococcus carnosus</i>	BaSTC2	Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Staphylococcus epidermidis</i>	1a, 2b, 3a, 4b, 5a, 6b, 7b, 8c, 9a, 10a, 11b, 12a & 13b	Can.J.Microbiol.1988.34:1358-1361
	41, 63, 118II, 138, 245, 336, 392 & 550	Res.Virol.1994.145:111-121
<i>Staphylococcus saprophyticus</i>	1154A, 1405, 1314, 1139 & 1259	Res.Virol.1990.141: 625-635 & Res.Virol.1994.145:111-121
<i>Staphylococcus sp.</i>	Phi 812, Phi 131, SK311 & U16	Virology 1998.246:241-252
<i>Streptococcus faecalis</i>	VD13	HER44 Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Streptococcus faecium</i>	PE1	Zentralbl.Bakteriol.1975.231:421-425
<i>Streptococcus oralis</i>	Cp-1** & Cp- 7**	FEMS Microbiol.Lett.1989.65:187-192

<i>Streptococcus pneumoniae</i>	Cp-1**	HER223	Felix d'Herelle Reference Centre, Quebec, Quebec
	Cp-1**, Cp-5**, Cp-7**, Cp-9**, ω -1 & ω -2		J.Virol.1981.40:551-559 & Eur.J.Biochem.1979.101:59-64 & Microbial Drug Resistance 1997.3:165-176
	HB-623 & HB-746		J.Virol.1990.64:5149-5155
	EJ-1**		J.Bacteriol.1992.174:5516-5525
	Dp-2 & Dp-4		J.Virol.1978.26:221-225
	Dp-1		Virology 1975.63:577-582
	ω -3 & ω -8		J.Virol.1976.19:659-667
	304		J.Bacteriol.1980.141:1298-1304
	HB-1, HB-2, HB-3**, HB-4, HB-5 & HB-6		J.Bacteriol.1979.138:618-624
<i>Streptococcus pyogenes</i>	T12**		Mol. Microbiology. 1997#23:719-728
	A-1	12202-B1	The American Type Culture Collection
	A-6	12203-B1	
	A-25	12204-B1	
<i>Streptococcus sp./Enterococcus</i>	Kjem	14918	
	1	HER 339	Felix d'Herelle Refrence Centre, Quebec, Quebec
	182	HER 80	
	VD1884	HER 323	
	1A	12169-B1	The American Type Culture Collection
	1B	12170-B1	
	NN	21597-B1	
	42	19948-B1	
	118	19951-B2	
	120	19952-B1	
<i>Veillonella rodentium</i>	N2		Antonie Van Leeuwenhoek 1989.56:263-271
<i>Vibrio cholerae</i>	Psi 92		Intervirology 1993.36:237-244
	VCB-1,2,3 & 4		J.Infection 1998.36:131
	CP-T1**		J.Virol.1984.51:163-169
	VSK		FEMS Microbiol.Lett.1996.145:17-22
	Phi138		J.Virol.1986.57:960-967
	Phi149		J.Virol.1985.140:217-223
	Fs-2**		Microbiology 1998.144:1901-1906

	e4 e5 X29 β κ 13 14 16 24 32 57		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Vibrio cholerae</i> (Cont'd)	138 145 149 163 N-4 S-5 S-20 M-4 D-10 I II III IV V	14100-B1 14100-B2 14100-B30 14100-B4 51352-B1 51352-B2 51352-B3 51352-B4 51352-B5 51352-b6 51352-B7 51352-B8 51352-B9 51352-B10	The American Type Culture Collection
<i>Vibrio costicola</i>	UTAK		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Vibrio eltor</i>	e ₄		J.Gen.Virol.1987.68:1411-1416
<i>Vibrio natrigens</i>	nt1, nt6		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Vibrio</i> <i>parahaemolyticus</i>	KVP40** VF33 VP1 ϕ 60 ϕ HAWI-5 ϕ PEL8C-1		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Vibrio sp.</i>	α 3a		Felix d'Herelle Reference Centre, Quebec, Quebec
	NN ph1	11985-B1 51582-B1	The American Type Culture Collection
	Phi149		J.Virol.1987.61:3999-4006
<i>Veillonella rodentium</i>	N2		Antonie V.Leeuwenhoek.1989.56:263-271

<i>Yersinia enterocolitica</i>	1 2 3 4 5 6 7 8 9 φYeO3-12		Felix d'Herelle Reference Centre, Quebec, Quebec
	I, IV & VIII		Zentralbl.Bakteriol.Mikrobiol.Hyg.1982.253:1 02
<i>Yersinia pestis</i>	R S Y	23208-B1 11593-B1 23053-B1	The American Type Culture Collection
	II		Zh.Mikrobiol.Epidemiol.Immunobiol.1990.11 :9
<i>Yersinia pseudotuberculosis</i>	PST**	23207-B1	The American Type Culture Collection
<i>Yersinia sp.</i>	RD2		Mol.Gen.Mikrobiol.Virusol.1990.8:18-21

xxxx)

Table 2

>Bacteriophage 77, complete genome sequence, 41708 nucleotides

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1      gatcaaaata cttggggaac ggtagggag taaacttcgc gataatttta aaaattcatg
61     tataaccccc ctcttataac cattttaagg caggtgatga aatggagatt atagtcgatg
121    aaaattttagt gcttaaagaa aaagaaaggc tacaagtatt atataaagac atacctagca
181    ataaattaaa agtagttgat ggtttaatta ttcaagcagc aaggctacgt gtaatgcttg
241    attacatgtg ggaagacata aaagaaaaag gtgattatga tttatttact caatctgaaa
301    aggcgcacc atagtaaagg gaaagaccag tagccaaact atttaatgct agagatgctg
361    catatcaaaa aataatcaaa caattatcgg atttattgcc cgaagagaaa gaagacacag
421    aaacgccatc tgatgattac ctatgattag taataaatac gttgatgaat atataaatTT
481    gtggaaacaa ggaaagataa ttttaataaa agaaagaatt gatctcttta attatctaca
541    aaaacatata tattcacgag atgatgtata ttttgatgaa cagaaaatcg aggattgtat
601    caaattttatt gaaaaatggg attttccaac attaccattt caaagggtta tcatagctaa
661    tatatttctt atagataaaa atacagatga agctttcttt acagaatttg ctattttcat
721    gggacgtgga ggcgggaaaa acggtctaata aagtgcattt agtgattttc tttctacgcc
781    cttacacgga gttaaagaat atcacatctc cattgttgct aatagtgaag atcaagcaaa
841    aacatcggtt gatgaaatca gaaccgtttt aatggataac aaacgaaata agacgggtaa
901    aacgcacaaa gctccttatg aagttagtaa agcaaaaata ataaaccgtg caactaaatc
961    ggttattcga tataacacat caaacacaaa aaccaagac ggtggacgtg aggggtgtgt
1021   tatttttgat gaaattcatt atttctttgg tcttgaaatg gtaaacgtca aacgtggtg
1081   attaggtaaa aagaaaaata gaagaacgtt ttatataagt actgatgggt ttgtagaga
1141   ggggttatatc gatgcaatga agcacaataa tgcaagtgtg ttaagtggca aggttaaaaa
1201   tagtagattg tttgcttttt attgtaagtt agacgatcca aaagaagttg atgacagaca
1261   gacgtgggaa aaggcgaacc caatgttaca taaaccgtta tcagaatacg ctaaaacact
1321   gctaagcacg attgaagaag aatataacga tttaccattc aaccgttcaa ataagcccga
1381   attcatgact aagcgaatga atttgcctga agttgacctt gaaaaagtaa tagcaccatg
1441   gaagaaataa ctacgcacta atagagagat accaaattta gataatcaaa tgtgtattgg
1501   tggtttagac tttgcaaaaca ttcgagattt tgcaagtgtg gggctattat tccgaaaaaa
1561   cgatgattac atttggttag gacattcgtt tgtaagacaa ggggttttgg atgatgtcaa
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Table 3

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2 77ORF006	3976..5196	49 77ORF053	37521..37757
3 77ORF007	21871..23076	50 77ORF054	22818..23060
4 77ORF008	2120..3307	51 77ORF055	17546..17788
5 77ORF009	31946..32803	52 77ORF058	18892..19122
6 77ORF010	26092..26889	53 77ORF059	34564..34785
7 77ORF011	24441..25208	54 77ORF064	29574..29795
8 77ORF012	29788..30576	55 77ORF065	28528..28746
9 77ORF013	33620..34399	56 77ORF066	27494..27703
10 77ORF014	27760..28512	57 77ORF069	38341..38547
11 77ORF015	3291..4028	58 77ORF070	36269..36475
12 77ORF016	32867..33610	59 77ORF071	40498..40701
13 77ORF017	23269..23982	60 77ORF072	38735..38938
14 77ORF018	31169..31840	61 77ORF073	30945..31148
15 77ORF019	39851..40501	62 77ORF074	38544..38738
16 77ORF020	6926..7570	63 77ORF075	13673..13870
17 77ORF021	37762..38304	64 77ORF077	25357..25605
18 77ORF022	30605..31156	65 77ORF079	29089..29280
19 77ORF023	26903..27346	66 77ORF080	35204..35389
20 77ORF024	10700..11140	67 77ORF085	24060..24242
21 77ORF025	9707..10147	68 77ORF092	39706..39876
22 77ORF026	40729..41145	69 77ORF094	32226..32393
23 77ORF027	6518..6925	70 77ORF096	13606..13773
24 77ORF028	34795..35199	71 77ORF098	7092..7256
25 77ORF029	6117..6521	72 77ORF102	29051..29212
26 77ORF030	36478..36879	73 77ORF104	34393..34551
27 77ORF031	39151..39546	74 77ORF109	18282..18434
28 77ORF032	33892..34266	75 77ORF112	39543..39692
29 77ORF033	5758..6120	76 77ORF117	27361..27501
30 77ORF034	7886..8236	77 77ORF118	38390..38530
31 77ORF035	19258..19560	78 77ORF120	36059..36199
32 77ORF036	36876..37223	79 77ORF124	33699..33833
33 77ORF037	102..446	80 77ORF128	14221..14355
34 77ORF038	34908..35219	81 77ORF130	15675..15806
35 77ORF039	37220..37528	82 77ORF133	8414..8542
36 77ORF040	41377..41676	83 77ORF140	13113..13235
37 77ORF041	35454..35753	84 77ORF147	7029..7148
38 77ORF042	5490..5774	85 77ORF149	30668..30787
39 77ORF043	29304..29564	86 77ORF151	31837..31953
40 77ORF044	18481..18768	87 77ORF155	30278..30391
41 77ORF045	5216..5500	88 77ORF157	4044..4157
42 77ORF046	25663..25935	89 77ORF167	20692..20799
43 77ORF047	11159..11425	90 77ORF175	35717..35821
44 77ORF048	28776..29039	91 77ORF176	6836..6940
45 77ORF049	36013..36255	92 77ORF178	35390..35491
46 77ORF050	35753..36007	93 77ORF179	8318..8419
47 77ORF051	38931..39167	94 77ORF182	29268..29564

Table 4

77ORF017 sequence

```

23982      atgacgcataatatagaaaaacgcattaataaattaaaaaacttct
1   M   T   H   N   I   E   K   R   I   N   K   L   K   T   S
23937      ggaaatccaaaattttaaaaagtttagattcagatattcactattta
16  G   N   P   K   F   K   K   L   D   S   D   I   H   Y   L
23892      ctcaagagatttgaaggtgaaaaaaaccataaagggtttttatcca
31  L   K   R   F   E   G   E   K   N   H   K   G   F   Y   P
23847      aagttttaacaaggagaaatagttttttagatttcggtataaac
46  K   F   K   Q   G   E   I   V   F   V   D   F   G   I   N
23802      gttaataaagaatttttctaattcacacttttgcaatagtgatgaat
61  V   N   K   E   F   S   N   S   H   F   A   I   V   M   N
23757      aaaaatgattctaatacggaggatatagtaaattgttattccctta
76  K   N   D   S   N   T   E   D   I   V   N   V   I   P   L
23712      tcctctaagaaaacaaaaagtttttaagatgaattttgatttg
91  S   S   K   E   N   K   K   Y   L   K   M   N   F   D   L
23667      aaatgggagttatttttaagattgtttttaaatttaatttagcgcg
106 K   W   E   Y   Y   L   R   L   F   L   N   L   I   S   A
23622      caaaataattcagctatattaaaagaagttttcgataaaaaatac
121 Q   N   N   S   A   I   L   K   E   V   F   D   K   K   Y
23577      caaaaaaacacacagaattcatcactaaagattattttattgaa
136 Q   K   N   N   T   E   F   I   T   K   D   Y   F   I   E
23532      tttatatctgatagtttagaaattgaaaataaattaaataaaaatt
151 F   I   S   D   S   L   E   I   E   N   K   L   N   K   I
23487      gacagaaacattaataacatagtatcagcaattgataaggtaaaa
166 D   R   N   I   N   N   I   V   S   A   I   D   K   V   K
23442      aaattaaaaggtaatagttacgcttgcataaattctttccagccg
181 K   L   K   G   N   S   Y   A   C   I   N   S   F   Q   P
23397      attagtaagtttcgcataagaaaagttttacccccaaaaaattaaa
196 I   S   K   F   R   I   R   K   V   L   P   Q   K   I   K
23352      aatccagtaatagattcttcggatattatgttactgataaataga
211 N   P   V   I   D   S   S   D   I   M   L   L   I   N   R
23307      attaataataatatattgcagatccctgatataagatga 23269
226 I   N   N   N   I   L   Q   I   P   D   I   R   *

```

Physico-chemical parameters of ORF 77ORF017

1 MTHNIEKRIN KLKTSGNPKF KKLDSDIHYL LKRFEGEKNH KGFYPKFKQG EIVFVDFGIN
 61 VNKEFSNSHF AIVMNKNSN TEDIVNVIPL SSKENKKYLK MNFDLKWEYY LRLFLNLISA
 121 QNNSAILKEV FDKKYQKNNT EFITKDYFIE FIDSLEIEN KLNKIDRNIN NIVSAIDKVK
 181 KLKGNYSYACI NSFQPISKFR IRKVLPOQIK NPVIDSSDIM LLINRINNNI LQIPDIR

Number of amino acids: 237
Average molecular weight (Daltons): 27887.38
Mean amino acid weight (Daltons): 117.67
Monoisotopic molecular weight (Daltons): 27869.83
Mean amino acid monoisotopic weight (Daltons): 117.59

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	5	2.11%	7.58%	Cys	C	1	0.42%	1.66%
Asp	D	14	5.91%	5.28%	Glu	E	13	5.49%	6.37%
Phe	F	16	6.75%	4.09%	Gly	G	6	2.53%	6.84%
His	H	4	1.69%	2.24%	Ile	I	29	12.24%	5.81%
Lys	K	33	13.92%	5.95%	Leu	L	19	8.02%	9.42%
Met	M	4	1.69%	2.37%	Asn	N	30	12.66%	4.45%
Pro	P	7	2.95%	4.9%	Gln	Q	6	2.53%	3.97%
Arg	R	8	3.38%	5.16%	Ser	S	17	7.17%	7.12%
Thr	T	5	2.11%	5.67%	Val	V	11	4.64%	6.58%
Trp	W	1	0.42%	1.23%	Tyr	Y	8	3.38%	3.18%

Number of acidic (negative) amino acids (ED): 27
 11.39%
Number of basic (positive) amino acids (KR): 41
 17.30%
Total charge (KRED): 68
 28.69%
Net charge (KR - ED): 14
 5.91%
Theoretical pI: 10.01
Total linear charge density: 0.30
Average hydrophobicity: -5.37
Ratio of hydrophilicity to hydrophobicity: 1.41
Percentage of hydrophilic amino acid: 57.81%
Percentage of hydrophobic amino acid: 42.19%
Ratio of %hydrophilic to %hydrophobic: 1.37

77ORF019 sequence

```
39851      atgaacgagcaaataataggaagcatatatacttttagcaggaggt
1   M   N   E   Q   I   I   G   S   I   Y   T   L   A   G   G
39896      gttgtgctttatttcagttaaagagatttttaggtattttacagat
16  V   V   L   Y   S   V   K   E   I   F   R   Y   F   T   D
39941      tctaacttacaacgtaaaaaaatcaatttagaacaaatatatccg
31   S   N   L   Q   R   K   K   I   N   L   E   Q   I   Y   P
39986      atatatttagattgtttttaaaaaggctaaaaagatgattggagct
46   I   Y   L   D   C   F   K   K   A   K   K   M   I   G   A
40031      tatattattccaacagaacagcatgaatttttagatttttttgat
61   Y   I   I   P   T   E   Q   H   E   F   L   D   F   F   D
40076      attgaagtctttaataatttagataagcaaagtaaaaaagcgtat
76   I   E   V   F   N   N   L   D   K   Q   S   K   K   A   Y
40121      gaaaatggtattggatttagacaaatgattaatttatcaaataga
91   E   N   V   I   G   F   R   Q   M   I   N   L   S   N   R
40166      gttaaggcaatggaagatttttaagatgagtttcaacaatgaattt
106  V   K   A   M   E   D   F   K   M   S   F   N   N   E   F
40211      agtacaaatcagattttttttaatccttcttttgttatggaaaca
121  S   T   N   Q   I   F   F   N   P   S   F   V   M   E   T
40256      attgctattataaatgaatatcaaaaagatatatcttattttaaaa
136  I   A   I   I   N   E   Y   Q   K   D   I   S   Y   L   K
40301      aatataattaataaaaatgaatgaaaatagagcttataatcatatt
151  N   I   I   N   K   M   N   E   N   R   A   Y   N   H   I
40346      gatagttttatcacttcagagtaccgacgaaaaataaacgattat
166  D   S   F   I   T   S   E   Y   R   R   K   I   N   D   Y
40391      aatctttatcttgataaatttgaagaacagtttagtcaaaaagttt
181  N   L   Y   L   D   K   F   E   E   Q   F   S   Q   K   F
40436      aaaataaacagaacttcgataaaagaaagaattattattaattta
196  K   I   N   R   T   S   I   K   E   R   I   I   I   N   L
40481      aacaagaggagattttaaatga 40501
211  N   K   R   R   F   K   *
```

Physico-chemical parameters of ORF 77ORF019

```

1      MNEQIIGSIY TLAGGVVLYS VKEIFRYFTD SNLQRKKINL EQIYPIYLDL FKKAKKMIGA
61     YIIPTEQHEF LDFFDIEVFN NLDKQSKKAY ENVIGFRQMI NLSNRVKAME DFKMSFNNEF
121    STNQIFFNPS FVMETIAIIN EYQKDISYLN NIINKMNENR AYNHIDSFIT SEYRRKINDY
181    NLYLDKFEEQ FSQKFKINRT SIKERIIINL NKRRFK

```

Number of amino acids: 216
Average molecular weight (Daltons): 26026.06
Mean amino acid weight (Daltons): 120.49
Monoisotopic molecular weight (Daltons): 26009.34
Mean amino acid monoisotopic weight (Daltons): 120.41

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	7	3.24%	7.58%	Cys	C	1	0.46%	1.66%
Asp	D	10	4.63%	5.28%	Glu	E	16	7.41%	6.37%
Phe	F	19	8.80%	4.09%	Gly	G	5	2.31%	6.84%
His	H	2	0.93%	2.24%	Ile	I	28	12.96%	5.81%
Lys	K	22	10.19%	5.95%	Leu	L	12	5.56%	9.42%
Met	M	7	3.24%	2.37%	Asn	N	23	10.65%	4.45%
Pro	P	3	1.39%	4.9%	Gln	Q	10	4.63%	3.97%
Arg	R	11	5.09%	5.16%	Ser	S	13	6.02%	7.12%
Thr	T	7	3.24%	5.67%	Val	V	7	3.24%	6.58%
Trp	W	0	0.00%	1.23%	Tyr	Y	13	6.02%	3.18%

Number of acidic (negative) amino acids (ED): 26
 12.04%
Number of basic (positive) amino acids (KR): 33
 15.28%
Total charge (KRED): 59
 27.31%
Net charge (KR - ED): 7
 3.24%
Theoretical pI: 9.52
Total linear charge density: 0.28
Average hydrophobicity: -4.84
Ratio of hydrophilicity to hydrophobicity: 1.37
Percentage of hydrophilic amino acid: 54.17%
Percentage of hydrophobic amino acid: 45.83%
Ratio of %hydrophilic to %hydrophobic: 1.18

77ORF043 sequence

```
29304      atgtattacgaaataggcgaaatcatacgcaaaaatattcatggt
1    M  Y  Y  E  I  G  E  I  I  R  K  N  I  H  V
29349      aacggattcgattttaagctattcattttaaaagggtcatatgggc
16   N  G  F  D  F  K  L  F  I  L  K  G  H  M  G
29394      atatcaatacaagttaaagatatgaacaacgtaccaattaaacat
31   I  S  I  Q  V  K  D  M  N  N  V  P  I  K  H
29439      gcttatgtcgtagatgagaatgacttagatatggcatcagactta
46   A  Y  V  V  D  E  N  D  L  D  M  A  S  D  L
29484      ttttaaccaagcaatagatgaatggattgaagagaacacagacgaa
61   F  N  Q  A  I  D  E  W  I  E  E  N  T  D  E
29529      caggacagactaattaacttagtcatgaaatggtag 29564
76   Q  D  R  L  I  N  L  V  M  K  W  *
```

Physico-chemical parameters of ORF 77ORF043

1 MYYEIGEIIIR KNIHVNGFDF KLFILKGHMG ISIQVKDMNN VPIKHAYVVD ENLDLMASDL
61 FNQAIDEWIE ENTDEQDRLI NLVMKW

Number of amino acids: 86
Average molecular weight (Daltons): 10186.68
Mean amino acid weight (Daltons): 118.45
Monoisotopic molecular weight (Daltons): 10180.02
Mean amino acid monoisotopic weight (Daltons): 118.37

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	3	3.49%	7.58%	Cys	C	0	0.00%	1.66%
Asp	D	9	10.47 %	5.28%	Glu	E	7	8.14%	6.37%
Phe	F	4	4.65%	4.09%	Gly	G	4	4.65%	6.84%
His	H	3	3.49%	2.24%	Ile	I	11	12.79 %	5.81%
Lys	K	6	6.98%	5.95%	Leu	L	6	6.98%	9.42%
Met	M	5	5.81%	2.37%	Asn	N	8	9.30%	4.45%
Pro	P	1	1.16%	4.9%	Gln	Q	3	3.49%	3.97%
Arg	R	2	2.33%	5.16%	Ser	S	2	2.33%	7.12%
Thr	T	1	1.16%	5.67%	Val	V	6	6.98%	6.58%
Trp	W	2	2.33%	1.23%	Tyr	Y	3	3.49%	3.18%

Number of acidic (negative) amino acids (ED): 16
18.60%
Number of basic (positive) amino acids (KR): 8
9.30%
Total charge (KRED): 24
27.91%
Net charge (KR - ED): -8
9.30%
Theoretical pI: 4.38
Total linear charge density: 0.30
Average hydrophobicity: -2.80
Ratio of hydrophilicity to hydrophobicity: 1.19
Percentage of hydrophilic amino acid: 48.84%
Percentage of hydrophobic amino acid: 51.16%
Ratio of %hydrophilic to %hydrophobic: 0.95

77ORF102 sequence

```
29051      atgagcaacattttataaaaagctacctagtagcagtattatgcttc
1      M  S  N  I  Y  K  S  Y  L  V  A  V  L  C  F
29096      acagtcttagcgattgtacttatgccgtttctataacttcactaca
16     T  V  L  A  I  V  L  M  P  F  L  Y  F  T  T
29141      gcatgggtcaattgcgggattcgcaagtatcgcaacattcatgtac
31     A  W  S  I  A  G  F  A  S  I  A  T  F  M  Y
29186      tacaaagaatgctttttcaaagaataa 29212
46     Y  K  E  C  F  F  K  E  *
```

Physico-chemical parameters of ORF 77ORF102

1 MSNIYKSYLV AVLCTVLAI VLMPLYFTT AWSIAGFASI ATFMYYKECF FKE

Number of amino acids: 53
Average molecular weight (Daltons): 6155.42
Mean amino acid weight (Daltons): 116.14
Monoisotopic molecular weight (Daltons): 6151.07
Mean amino acid monoisotopic weight (Daltons): 116.06

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	6	11.32 %	7.58%	Cys	C	2	3.77 %	1.66%
Asp	D	0	0.00%	5.28%	Glu	E	2	3.77 %	6.37%
Phe	F	7	13.21 %	4.09%	Gly	G	1	1.89 %	6.84%
His	H	0	0.00%	2.24%	Ile	I	4	7.55 %	5.81%
Lys	K	3	5.66%	5.95%	Leu	L	5	9.43 %	9.42%
Met	M	3	5.66%	2.37%	Asn	N	1	1.89 %	4.45%
Pro	P	1	1.89%	4.9%	Gln	Q	0	0.00 %	3.97%
Arg	R	0	0.00%	5.16%	Ser	S	4	7.55 %	7.12%
Thr	T	4	7.55%	5.67%	Val	V	4	7.55 %	6.58%
Trp	W	1	1.89%	1.23%	Tyr	Y	5	9.43 %	3.18%

Number of acidic (negative) amino acids (ED): 2
 3.77%
Number of basic (positive) amino acids (KR): 3
 5.66%
Total charge (KRED): 5
 9.43%
Net charge (KR - ED): 1
 1.89%
Theoretical pI: 8.18
Total linear charge density: 0.13
Average hydrophobicity: 10.81
Ratio of hydrophilicity to hydrophobicity: 0.40
Percentage of hydrophilic amino acid: 28.30%
Percentage of hydrophobic amino acid: 71.70%

Ratio of %hydrophilic to %hydrophobic:

0.39

77ORF104 sequence

```
34393      atggtaaccaaagaattttttaaaaactaaacttgagtgttcagat
1      M  V  T  K  E  F  L  K  T  K  L  E  C  S  D
34438      atgtacgctcagaaaactcatagatgaggcacagggcgatgaaaat
16     M  Y  A  Q  K  L  I  D  E  A  Q  G  D  E  N
34483      aggttgtagacacctatctatccaaaaacttgcagaacgtcataca
31     R  L  Y  D  L  F  I  Q  K  L  A  E  R  H  T
34528      cgccccgctatcgtcgaatattaa 34551
46     R  P  A  I  V  E  Y  *
```


Physico-chemical parameters of ORF 77ORF104

1 MVTKEFLKTK LECDMYAQK LIDEAQGDEN RLYDLFIQKL AERHTRPAIV EY

Number of amino acids: 52
 Average molecular weight (Daltons): 6193.13
 Mean amino acid weight (Daltons): 119.10
 Monoisotopic molecular weight (Daltons): 6189.12
 Mean amino acid monoisotopic weight (Daltons): 119.02

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	4	7.69 %	7.58%	Cys	C	1	1.92%	1.66%
Asp	D	4	7.69 %	5.28%	Glu	E	6	11.54 %	6.37%
Phe	F	2	3.85 %	4.09%	Gly	G	1	1.92%	6.84%
His	H	1	1.92 %	2.24%	Ile	I	3	5.77%	5.81%
Lys	K	5	9.62 %	5.95%	Leu	L	6	11.54 %	9.42%
Met	M	2	3.85 %	2.37%	Asn	N	1	1.92%	4.45%
Pro	P	1	1.92 %	4.9%	Gln	Q	3	5.77%	3.97%
Arg	R	3	5.77 %	5.16%	Ser	S	1	1.92%	7.12%
Thr	T	3	5.77 %	5.67%	Val	V	2	3.85%	6.58%
Trp	W	0	0.00 %	1.23%	Tyr	Y	3	5.77%	3.18%

Number of acidic (negative) amino acids (ED): 10
 19.23%
 Number of basic (positive) amino acids (KR): 8
 15.38%
 Total charge (KRED): 18
 34.62%
 Net charge (KR - ED): -2
 3.85%
 Theoretical pI: 5.03
 Total linear charge density: 0.38
 Average hydrophobicity: -5.81
 Ratio of hydrophilicity to hydrophobicity: 1.47
 Percentage of hydrophilic amino acid: 53.85%
 Percentage of hydrophobic amino acid: 46.15%

Ratio of %hydrophilic to %hydrophobic:

1.17

77ORF182 sequence

```
29268      atgttcaatataaaaacgaaaaacggaggaagtcaagatgtattac
1      M  F  N  I  K  R  K  T  E  E  V  K  M  Y  Y
29313      gaaataggcgaaatcatatcgcaaaaatattcatgttaacggattc
16     E  I  G  E  I  I  R  K  N  I  H  V  N  G  F
29358      gattttaagctattcatttttaaagggtcatatgggcatatcaata
31     D  F  K  L  F  I  L  K  G  H  M  G  I  S  I
29403      caagttaaagatatgaacaacgtaccaattaaacatgcttatgtc
46     Q  V  K  D  M  N  N  V  P  I  K  H  A  Y  V
29448      gtagatgagaatgacttagatatggcatcagacttatttaaccaa
61     V  D  E  N  D  L  D  M  A  S  D  L  F  N  Q
29493      gcaatagatgaatggattgaagagaacacagacgaacaggacaga
76     A  I  D  E  W  I  E  E  N  T  D  E  Q  D  R
29538      ctaattaacttagtcatgaaatggtag 29564
91     L  I  N  L  V  M  K  W  *
```

Physico-chemical parameters of ORF 77ORF182

1 MFNIKRKTEE VKMYEIGEI IRKNIHVNGF DFKLFILKGH MGISIQVKDM NNVPIKHAYV
 61 VDENDLDMAS DLFNQAIDEW IEENTDEQDR LINLVMKW

Number of amino acids: 98
 Average molecular weight (Daltons): 11691.50
 Mean amino acid weight (Daltons): 119.30
 Monoisotopic molecular weight (Daltons): 11683.84
 Mean amino acid monoisotopic weight (Daltons): 119.22

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	3	3.06 %	7.58%	Cys	C	0	0.00%	1.66%
Asp	D	9	9.18 %	5.28%	Glu	E	9	9.18%	6.37%
Phe	F	5	5.10 %	4.09%	Gly	G	4	4.08%	6.84%
His	H	3	3.06 %	2.24%	Ile	I	12	12.24 %	5.81%
Lys	K	9	9.18 %	5.95%	Leu	L	6	6.12%	9.42%
Met	M	6	6.12 %	2.37%	Asn	N	9	9.18%	4.45%
Pro	P	1	1.02 %	4.9%	Gln	Q	3	3.06%	3.97%
Arg	R	3	3.06 %	5.16%	Ser	S	2	2.04%	7.12%
Thr	T	2	2.04 %	5.67%	Val	V	7	7.14%	6.58%
Trp	W	2	2.04 %	1.23%	Tyr	Y	3	3.06%	3.18%

Number of acidic (negative) amino acids (ED): 18
 18.37%
 Number of basic (positive) amino acids (KR): 12
 12.24%
 Total charge (KRED): 30
 30.61%
 Net charge (KR - ED): -6
 6.12%
 Theoretical pI: 4.76
 Total linear charge density: 0.33
 Average hydrophobicity: -3.89
 Ratio of hydrophilicity to hydrophobicity: 1.28

167

Percentage of hydrophilic amino acid:	51.02%
Percentage of hydrophobic amino acid:	48.98%
Ratio of %hydrophilic to %hydrophobic:	1.04

Table 5

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100017|lan|77ORF017 Phage 77 ORF |23269-23982|-3
(237 letters)

Database: nr

393,678 sequences; 120,452,765 total letters

Sequences producing significant alignments:	Score (bits)	E Value
gi 4493986 emb CAB39045.1 (AL034559) predicted using hexExon; ...	41	0.010
gi 730607 sp P23250 RPI1_YEAST NEGATIVE RAS PROTEIN REGULATOR P...	38	0.053
gi 3097044 emb CAA75299 (Y15035) K1R [Cowpox virus]	38	0.090
gi 2146245 pir S73794 hypothetical protein H91_orf180 - Mycopl...	38	0.090
gi 83910 pir S04682 ribosomal protein var1 - yeast (Candida gl...	37	0.15
gi 133135 sp P21358 RMAR_CANGA MITOCHONDRIAL RIBOSOMAL PROTEIN ...	37	0.15
gi 2128843 pir H64475 hypothetical protein MJ1409 - Methanococ...	36	0.20
gi 5107017 gb AAD39926.1 AF126285_2 (AF126285) RNA polymerase {...	36	0.35
gi 2146210 pir S73342 hypothetical protein E07_orf166 - Mycopl...	35	0.60

Database: swissprot

79,449 sequences; 28,874,452 total letters

Sequences producing significant alignments:	Score (bits)	E Value
sp P23250 RPI1_YEAST NEGATIVE RAS PROTEIN REGULATOR PROTEIN.	38	0.014
sp P21358 RMAR_CANGA MITOCHONDRIAL RIBOSOMAL PROTEIN VAR1.	37	0.040
sp Q21444 LDLC_CAEEL LDLC PROTEIN HOMOLOG.	34	0.35
sp P27240 RFAY_ECOLI LIPOPOLYSACCHARIDE CORE BIOSYNTHESIS PROT.	33	0.46
sp P53192 YGCO_YEAST HYPOTHETICAL 27.1 KD PROTEIN IN ALK1-CKB1.	33	0.60
sp P32908 SMC1_YEAST CHROMOSOME SEGREGATION PROTEIN SMC1 (DA-B.	33	0.60
sp P54683 TAGB_DICDI PRESTALK-SPECIFIC PROTEIN TAGB PRECURSOR .	32	0.78
sp Q03100 CYAA_DICDI ADENYLATE CYCLASE, AGGREGATION SPECIFIC (.	32	0.78

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BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100019|lan|77ORF019 Phage 77 ORF|39851-40501|2
(216 letters)

Database: nr
373,355 sequences; 114,214,446 total letters

Sequences producing significant alignments:	Score (bits)	E Value
gi 3341966 dbj BAA31932 (AB009866) orf 59 [bacteriophage phi PVL]	437	e-122
gi 2689911 (AE000792) B. burgdorferi predicted coding region BB...	38	0.058
gi 1171589 emb CAA64574 (X95275) frameshift [Plasmodium falcip...	37	0.10
gi 4493986 emb CAB39045.1 (AL034559) predicted using hexExon; ...	36	0.23
gi 141257 sp P18019 YPI9_CLOPE HYPOTHETICAL 14.5 KD PROTEIN (OR...	36	0.29
gi 133412 sp P27059 RPOB_ASTLO DNA-DIRECTED RNA POLYMERASE BETA...	35	0.51
gi 3122231 sp Q58851 HISX_METJA HISTIDINOL DEHYDROGENASE (HDH) ...	35	0.51
gi 3649757 emb CAB11106.1 (Z98547) predicted using hexExon; MA...	34	0.66
gi 2688313 (AE001146) sensory transduction histidine kinase, pu...	34	0.87

Database: swissprot
79,449 sequences; 28,874,452 total letters

Sequences producing significant alignments:	Score (bits)	E Value
sp P18019 YPI9_CLOPE HYPOTHETICAL 14.5 KD PROTEIN (ORF9).	36	0.079
sp Q58851 HISX_METJA HISTIDINOL DEHYDROGENASE (EC 1.1.1.23) (H.	35	0.14
sp P27059 RPOB_ASTLO DNA-DIRECTED RNA POLYMERASE BETA CHAIN (E.	35	0.14
sp Q02224 CENE_HUMAN CENTROMERIC PROTEIN E (CENP-E PROTEIN).	34	0.31
sp P04931 ARP_PLAFA ASPARAGINE-RICH PROTEIN (AG319) (ARP) (FRA..	33	0.53
sp P18011 IPAB_SHIFL 62 KD MEMBRANE ANTIGEN.	32	0.69
sp P18709 VTA2_XENLA VITELLOGENIN A2 PRECURSOR (VTG A2) [CONTA..	32	0.90
sp Q64409 CP3H_CAVPO CYTOCHROME P450 3A17 (EC 1.14.14.1) (CYPI..	32	0.90
sp P21358 RMAR_CANGA MITOCHONDRIAL RIBOSOMAL PROTEIN VAR1.	32	0.90
sp Q03945 IPAB_SHIDY 62 KD MEMBRANE ANTIGEN.	32	1.2

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BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100043|lan|77ORF043 Phage 77 ORF|29304-29564|3
(86 letters)

Database: nr
373,355 sequences; 114,214,446 total letters

Sequences producing significant alignments:	Score (bits)	E Value
gi 3341947 dbj BAA31913 (AB009866) orf 39 (bacteriophage phi PVL)	182	6e-46
gi 744518 prf 2014422A FKBP-rapamycin-associated protein [Homo...	32	0.84
gi 1169736 sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN...	32	0.84
gi 1169735 sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTE...	32	0.84
gi 3282239 (U88966) rapamycin associated protein FRAP2 [Homo sa...	32	0.84
gi 3875402 emb CAA98122 (Z73906) cDNA EST EMBL:D64544 comes fr...	31	2.5
gi 1084792 pir S54091 hypothetical protein YPR070w - yeast (Sa...	30	4.2

Database: swissprot
79,449 sequences; 28,874,452 total letters

Sequences producing significant alignments:	Score (bits)	E Value
sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) .	32	0.24
sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R.	32	0.24
sp P34554 YNP1_CAEEL HYPOTHETICAL 42.2 KD PROTEIN T05G5.1 IN C.	28	3.5
sp Q24118 LIO_DROME LINOTTE PROTEIN.	28	3.5
sp P80034 ACH2_BOMMO ANTICHYMOTRYPSIN II (ACHY-II).	28	3.5
sp P22922 ALAT_BOMMO ANTITRYPSIN PRECURSOR (AT).	28	3.5
sp Q44363 TRAA_AGRT6 CONJUGAL TRANSFER PROTEIN TRAA.	28	3.5
sp P38255 YBU5_YEAST HYPOTHETICAL 51.3 KD PROTEIN IN PHO5-VPS1.	27	6.0
sp P55822 SH3B_HUMAN SH3BGR PROTEIN (21-GLUTAMIC ACID-RICH PRO.	27	7.9
sp Q58482 YA82_METJA HYPOTHETICAL PROTEIN MJ1082.	27	7.9
sp P34252 YKK8_YEAST HYPOTHETICAL 52.3 KD PROTEIN IN HAP4-AAT1.	27	7.9

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BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100102|lan|77ORF102 Phage 77 ORF|29051-29212|2
(53 letters)

Database: nr
373,355 sequences; 114,214,446 total letters

	Score (bits)	E Value
Sequences producing significant alignments:		
gi 3341946 dbj BAA31912 (AB009866) orf 38 [bacteriophage phi PVL]	96	3e-20
gi 4325288 gb AAD17315 (AF123593) voltage-dependent sodium cha...	28	7.1
gi 2649684 (AE001040) A. fulgidus predicted coding region AF092...	28	9.3

Database: swissprot
79,449 sequences; 28,874,452 total letters

	Score (bits)	E Value
Sequences producing significant alignments:		
sp P42087 HUTM_BACSU PUTATIVE HISTIDINE PERMEASE.	26	7.1
sp P04775 CIN2_RAT SODIUM CHANNEL PROTEIN, BRAIN II ALPHA SUBU...	26	9.2
sp P42619 YQJF_ECOLI HYPOTHETICAL 17.2 KD PROTEIN IN EXUR-TDCC...	26	9.2

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BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100104|lan|77ORF104 Phage 77 ORF|34393-34551|1
(52 letters)

Database: nr
373,355 sequences; 114,214,446 total letters

Sequences producing significant alignments:	Score (bits)	E Value
gi 2315523 (AF016452) similar to the leucine-rich domains found...	29	4.2
gi 4377168 gb AAD18990 (AE001666) CT711 hypothetical protein [...]	29	5.4
gi 3882171 dbj BAA34445 (AB018268) KIAA0725 protein [Homo sapi...]	28	9.3

Database: swissprot
79,449 sequences; 28,874,452 total letters

Sequences producing significant alignments:	Score (bits)	E Value
sp P04879 RRPP_VSVIG RNA POLYMERASE ALPHA SUBUNIT (EC 2.7.7.48.	27	5.4
sp P04880 RRPP_VSVIM RNA POLYMERASE ALPHA SUBUNIT (EC 2.7.7.48.	27	5.4
sp Q13946 CN7A_HUMAN HIGH-AFFINITY CAMP-SPECIFIC 3',5'-CYCLIC .	26	7.1
sp P35381 ATPA_DROME ATP SYNTHASE ALPHA CHAIN, MITOCHONDRIAL P.	26	9.3
sp P54659 MVPB_DICDI MAJOR VAULT PROTEIN BETA (MVP-BETA) .	26	9.3
sp P40397 YHXC_BACSU HYPOTHETICAL OXIDOREDUCTASE IN APRE-COMK .	26	9.3

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BLASTP 2.0.8 [Jan-05-1999]

Query= sid|122748|lan|77ORF182 Phage 77 ORF|29268-29564|3
(98 letters)

Database: nr
393,678 sequences; 120,452,765 total letters

Sequences producing significant alignments:	Score (bits)	E Value
gi 3341947 dbj BAA31913.1 (AB009866) orf 39 [bacteriophage phi..	182	8e-46
gi 1084792 pir S54091 hypothetical protein YPR070w - yeast (Sa..	35	0.13
gi 1169736 sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN..	32	1.1
gi 744518 prf 2014422A FKBP-rapamycin-associated protein [Homo..	32	1.1
gi 5051381 emb CAB44736.1 (AL049653) dJ647M16.2 (FK506 binding..	32	1.1
gi 4826730 ref NP_004949.1 pFRAP1 FK506 binding protein 12-rap..	32	1.1
gi 3282239 (U88966) rapamycin associated protein FRAP2 [Homo sa..	32	1.1

Database: swissprot
79,909 sequences; 29,054,478 total letters

Sequences producing significant alignments:	Score (bits)	E Value
sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) .	32	0.29
sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R..	32	0.29
sp P40557 YIA5_YEAST PUTATIVE DISULFIDE ISOMERASE YIL005W PREC..	29	3.3
sp Q24118 LIO_DROME LINOTTE PROTEIN.	28	4.4
sp Q44363 TRAA_AGRT6 CONJUGAL TRANSFER PROTEIN TRAA.	28	4.4
sp P80034 ACH2_BOMMO ANTICHYMOTRYPSIN II (ACHY-II).	28	4.4
sp P34554 YNP1_CAEEL HYPOTHETICAL 42.2 KD PROTEIN T0SG5.1 IN C..	28	4.4
sp P22922 ALAT_BOMMO ANTITRYPSIN PRECURSOR (AT).	28	4.4

Table 6

1st position (5' end)	2nd position				3rd position (3' end)
	U	C	A	G	
U	Phe	Ser	Tyr	Cys	U
	Phe	Ser	Tyr	Cys	C
	Leu	Ser	Stop	Stop	A
	Leu	Ser	Stop	Trp	G
C	Leu	Pro	His	Arg	U
	Leu	Pro	His	Arg	C
	Leu	Pro	Gln	Arg	A
	Leu	Pro	Gln	Arg	G
A	Ile	Thr	Asn	Ser	U
	Ile	Thr	Asn	Ser	C
	Ile	Thr	Lys	Arg	A
	Met	Thr	Lys	Arg	G
G	Val	Ala	Asp	Gly	U
	Val	Ala	Asp	Gly	C
	Val	Ala	Glu	Gly	A
	Val	Ala	Glu	Gly	G

Table 7

Bacteriophage 3A, complete genome sequence

1	caaacgctag	caacgcggat	aaatttttca	tgaaggggg	tctttatatg	aagttaacaa	aaaaacagct
71	aaaagaatat	atagaagatt	acaaaaaatc	tgatgacata	ttaattaatt	tgatatataga	aacatatgaa
141	ttttattgtc	gggttaagaga	tgaacttaaa	aatagtgatt	taatgataga	gcatacaaac	aaggctgggtg
211	cgagcaatat	tattaagaat	ccattaaagca	tagaactgac	aaaaacagct	caaacactaa	ataacttact
281	caagtcctatg	gggttaactg	cagcacaaag	aaaaaagata	gttcaagaag	aaggtgggatt	cgggtgactat
351	taaagtttta	aatgaacctt	cacaaaaact	attaacaaca	tggtatgcag	agcaagtcac	tcaagggaata
421	ataaaaacaa	gcaaatatgt	tagaaaaagaa	tgtgagagac	atcttagata	tctagaaaaat	ggaggttaaat
491	gggtatttga	tgaagaatta	gcgcacgtc	ctattcgatt	tatagaaaaag	ttttgtaaac	cttccaaagg
561	atctaaacgt	caacttgat	tacagccatg	gcaacatttt	attatcgga	gtttgtttgg	ttgggttcat
631	aaagaaacaa	aactgcgcag	gtttaaagaa	gctttgatat	ttatggggcg	aaaaaatggt	aaaacaacca
701	ctatttctgg	gggtgctaac	tatgctgtat	cacaagatgg	agaaaaatggt	gcagaaattc	atttgttagc
771	aaacgttaag	aaacaagcta	ggatttctatt	tgatgaatct	aaggcgatga	ttaagctag	cccaagctt
841	gatcaaaatt	tcagaacatt	aagagatgaa	atccattatg	acgcaacgat	atcaaaaatt	atgccccaaag
911	catcagatag	cgataagtta	gatggattga	atacacacat	ggggattttt	gatgaaattc	atgaatttaa
981	agactataaa	ttgatttctag	ttataaaaaa	ctcaagagct	gcaagggtac	aacctcttct	catctacatt
1051	acgacagcag	gggtatcaatt	agatgggtcca	cttgttgata	tggtagaagc	gggaagagac	accttagatc
1121	aaatcataga	agacgaaaga	actttttatt	atttagcatc	tttggatgat	gaogatgata	ttaatgattc
1191	gtcgaaactg	ataaaagcaa	atcccaactt	aggtgtctct	ataaaattag	atgagatgaa	agaagagtgg
1261	gaaaaagcta	agagaacacc	agctgaaact	ggagatttta	taaccaaaag	gtttaatatc	tttgcttaata
1331	atgacgagat	gagttttatt	gattacccaa	cactccaaa	aaataatgaa	attgtttctt	tagaagagct
1401	ggaagggcaga	ccgtgcacga	ttgggtatga	tttatcagaa	acagaggact	ttacagccgc	gtgtgctact
1471	tttgcgttag	ataatggtaa	agttgcagtt	ttatcgcat	catggattcc	taagcacaaa	gttgaattatt
1541	ctaacgaaaa	aataccctat	agagaatggg	aagaagatgg	cttattaaca	gtgcaagata	agccttatat
1611	tgactaccaa	gatgttttaa	atgggataat	taagatgaat	gagcattatg	tagtagaaaa	aattacttat
1681	gatagagcga	acgcattcaa	actaaatcaa	gagttaaaaa	attacggggt	tgaaacggaa	gaaacaagac
1751	aaggagcttt	gactttgagc	ctcgtatga	aggatttaaa	agaaatgttt	ttagatggga	aaataatatt
1821	taataataat	gcttttaata	aatgggtatat	caataatggt	cagttgaaac	tagacagaaa	cggaaactgg
1891	ttgcgctcta	agcaaaagcag	atatcgtaaa	atagatgggt	ttgcagcatt	tttaaacaca	tatacagata
1961	ttatgaataa	agttgtttct	gatagtgggt	aagggaacat	agagtttatt	agtattaaag	acataatgcg
2031	ttaaggagggt	gaatgttatc	gcaaaagaga	atattgtcac	acgcataaag	aaaaaattga	tagacaattg
2101	gattgatcag	tcaacttcta	agctttatga	ctttagccca	tggaaaaaata	gatctttttg	gggtgtaatt
2171	aataatcacg	ttgaaactaa	tgaaacgata	ttttcagcta	ttacaagatt	atctaattcg	atggctagtt
2241	tgcccttgaa	aatgtatgaa	gattataaag	tagttaatac	agaagtatct	gatttactta	cagttgtcacc
2311	gaataaattc	ctgagcagtt	ttgattttat	taatacaatt	gaaacaatca	gaaatgaaaa	aggtaatgca
2381	tatgtgtcaa	ttgaacgaga	catctatcat	caaccatcaa	agcttttctt	attaaatcca	gatgttggtg
2451	aaatgttaat	tgaaaaccaa	tcacgtgaac	tttattatc	cattcatgct	gcaactggaa	ataaattgat
2521	tggtcataat	atggacatgt	tgcattttaa	acacatcggt	gcattctaata	tggtgcaagg	cattagtccg
2591	attgatgtgt	tgaagaatac	aactgatttt	gataatgcag	taagaacctt	taactctaca	gaaatgcaaa
2661	aacctgatcc	tttcatgctt	aaatatgggt	ccaatgtagg	taagaaaaaa	aggcagcaag	tgtagaaga
2731	tttcaaacag	tactatgaag	aaaacggtgg	aatattatc	caagagccgt	gtgttgaaat	cgaacggtta
2801	cctaaaaaat	atgctctcta	agatatagtg	gcaagcgaga	atttaacaag	agaaagagta	gctaacgttt
2871	ttcaattgcc	ctcagtatcc	ttaaatgcaa	gatcaaatac	aaatttccgc	aaaaatgaag	agttaaacag
2941	attttacttg	cagatatact	tattgccaat	cgtaaacacg	tatgaagaag	aattttaatcg	gaaactactt
3011	actaaaaacg	acagagaaaa	aaataggtat	tttaaaattta	acgttaaaac	ttatttaagg	gctgatagtg
3081	caacacagc	agaagtgtac	tttaagcag	ttcgtagtgg	ttactacact	ataaatgaca	ttagagagtg
3151	ggaagattta	ccaccagttg	aaggtggaga	taagccgcta	ataagcgggt	atttataacc	aattgcacag
3221	ccacttgaa	taagaaaaat	tttgaaagg	ggtgataaaa	atgtcaatga	aagctaagta	ttttcaaatg
3291	aaaagaaaa	caaaaagtaa	aggtgaaata	tttatttatg	gtgatattgt	aagtgataaa	tggtttgaaa
3361	gtgatgtaac	tgctacagat	ttcaaaaaata	aactagatga	actaggagac	atcagtgaat	tagatgttca
3431	tataaattca	tctggaggca	gtgtatttga	agggcatgca	atatacaata	tgctaaaaat	gcatcctgca
3501	aaaatttaata	tctatgtcga	tgcttagcgc	gcatcaattg	ctagtgttat	cgctatgagt	ggtgacacta
3571	tttttatgca	caaaaatagt	tttttaata	ttcataattc	atggggttatg	actgtaggta	atgcagaaga
3641	gttaagaaag	acagcggatt	tacttgaaaa	aacagatgct	gttagtaatt	cagcttattt	agataaagca
3711	aaagatttag	atcaagaaca	cttaaaacag	atgttagatg	cagaaacttg	gcttactgca	gaagaagcct
3781	tgctcttcgg	cttgatagat	gaaatttttag	gagctaata	aataactgct	agtatctcta	aagagcaata
3851	taagcgtttc	gagaaactcc	cagaagattt	aaagaaagat	gtagacaaaa	tcactaaaat	cgatgatgta
3921	gatacgtttg	aattgggtga	aacacctaata	gaaagtatgt	cactagaaga	aaaagaaaaa	agagaaaaaa
3991	ttaaacgcga	atgcgaaatt	ttaaaaatga	caatgagtta	ttaggaggaa	atgaaatgcc	gacattatat
4061	gaaatttaaac	aatccttagg	tatgatttga	caacaattaa	aaaataaaaa	tgatgaattg	agtcagaaaag
4131	caacagaccc	aatatttgat	atggaagaca	tcaacaactc	agaaacagaa	aaagcaggct	tacaacaaag
4201	attttaacatt	gttgaaagac	agtaaaaaa	cattgaagaa	aaagaaaaag	cgaaagttaa	agacacagga
4271	gaagctttac	aatcttttaa	tgatcatgag	aagctgggtta	aagctaaggc	agagttttat	cgtaacgcga
4341	ttttaccaaa	tgaatttgaa	aaaccttcaa	tggaaggcaca	acgtttatta	cacgctttac	caacaggtaa
4411	tgattcaggt	ggtgataagc	ctttaccaaa	aacactttct	aaagaaattg	tttcagaacc	atttgctaaa
4481	aaccaattac	gtcaaaaagc	tcgtctaact	aacattaaag	gtttagagat	tccaagagtt	tcataactt
4551	tagacgatga	tgacttcatt	acagatgtga	aaacagcaaa	agaattaaaa	ttaaaagggtg	atacagttaa
4621	attcactact	ataaattca	aagtatttgc	tgcaatttca	gatactgtaa	ttcatggatc	agatgtagat
4691	ttagttaaact	gggttgaaaa	cgactacaaa	tcagggtctag	cagctaaaga	acgtaaaagat	gccttagcag

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4761 taagtcctaa atctggatta gatcacatgt cattttacaa tggatctgtt aaagaagtgt agggagcaga
4831 catgtatgat gctattatta acgcttttagc agattttacat gaagattacc gtgataacgc aacaatttat
4901 atcgcatatgt cggattattgt caaaattatt agtggtctttt caaatggaac aacaaatttc tttgacacac
4971 cagcagaaaa agtatttggc aaaccagtag tatttacaga tgcagcagtt aaacctattg tgggagattt
5041 caattattttt ggaattaaact atgatggaaac aactttatgac actgataaag atgttaaaaa aggcgaatat
5111 ttgtttgtat taactgcatg gtatgatcag caacgtacat tagacagtgct attcagaatt gcaaaagcaa
5181 aaagaaaatc aggttcatta cccagctaag ccccaaaagg ttaatgtaac agctaaggct aaatcagctg
5251 taatatcagc cgaatagggg tgatgaaatg agtttagaag aaattaaatt gtggttgaga attgactata
5321 atttcgaaaa tgatttaatt gaaggtctca ttcaatcggc taagtctgaa ttactattaa gtggggtccc
5391 agattatgac aaagatgact tggaaatccc gcttttttgt acagcgatta gatatatcat tgcaagagat
5461 tatgaaagtc gtgggtactc aaatgaccaa tctagaagca aggtttttta tgaaaaggga ttgcaaaaaa
5531 tgattctgaa attaaaaaag tggtaggtga tttttaaatg gaatttaattg aatttaaaaga tccgcgcatt
5601 ttttttcaat atgtaataaa agggccgtat ccagatgaag aggaaaaaat gaagttgtat agttgctttt
5671 gaaaaatata taatccctct atgaaagata gagaaatttt aaaagcgact gaatcaaatg caggactaac
5741 cataattatg aggtcttcta aaattgaata tctaccacaa acaaatcact tagttaaaat tgacagaggc
5811 ttatttccg ataaattatt caacattaaa gaaataagaa ttgatcaccc agatattggc tataatacac
5881 tgggtttttt agaaaaatga gtgtagaat taaggggata cctgaagtgt tgaagaaatt agaatcggta
5951 tacggtaaac aatcaatgca agctaagagt gatagagctt taaatgaagc atctgaattt tttataaagg
6021 ctttaagaa agaattcgag agttttaaag atacgggtgc tagcatagaa gaaatgacta aatctaagcc
6091 ttatacaaaa gttaggaagtc aagaaagagc tgttttaatt gaatgggtag gccctatgaa tcgcaaaaac
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Table 8

Bacteriophage 3A ORFs list

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100382	3AORF004	3	17457..19370	637	gctattttattagaaaggaaggtgc	att	taa
100383	3AORF005	1	334..2034	566	agaaaaagatagttcaagaagaag	gtg	taa
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100386	3AORF008	3	22176..23630	484	aatgatttagggtaggtgttgacca	atg	tga
100387	3AORF009	1	40726..42093	455	gtaaatacttttataagaatggtag	gtg	taa
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100389	3AORF011	2	2039..3277	412	attaaagacataatgcggttaaggag	gtg	taa
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100653	3AORF275	3	39882..39995	37	gatatgttaccacaggaatgtag	att	taa
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100657	3AORF279	-2	17535..17648	37	tttgtaaagatttgtttactgctgc	ttg	taa
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100659	3AORF281	-2	759..872	37	ttttgatatcggtgctcataatgg	att	tga
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100717	3AORF339	3	27348..27449	33	atatctaattaaataagcgcaactta	att	tga
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100722	3AORF344	-1	7297..7398	33	ctgctgaaactgttgcagattttga	att	tga
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Table 9

Bacteriophage 96, complete genome sequence

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 36261 aagatgaaaa gtcacaggtg taatgtctga ggctttttaa ttaacacaa agtaggtggc gtaatgtttg
 36331 gatttaccaa acggcacgaa catgaatggc gaattagaag attagaagag aatgataaaa caatgcttaa
 36401 cactctcaat gagattaaat taggtcaaaa aactcaagag caagttaaca ttaaattaga taaaacttta
 36471 gtgctatcc agaggggaaag acagatagac gaaaaaata agaaagaaaa cgacaaaaat atacggcata
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 38921 aacacgctgg ttatgttcgc tgaagctgt ttttagagaa agaagaagaa aaaatacatg tcgcaaaact
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 39691 caagcggata atttatatca actaaaagc gcactcaac cgacgggtta aatttggaca ggaacagaaa
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Table 10

Bacteriophage 96 ORFs list

SID	LAN	FRA	POS	a.a.	RBS sequence	STA	STO
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100735	96ORF003	1	30109..31995	628	ttatatttttagataaggagtagcct	atg	taa
100736	96ORF004	1	36760..38634	624	attttgattgaaatgaggtgcatac	atg	taa
100737	96ORF005	3	33903..35729	608	gtttattcgaaggaaaggtggttga	ata	taa
100738	96ORF006	2	40589..42043	484	aatgatttagggtagggtgttgacca	atg	tag
100739	96ORF007	1	18652..20091	479	tatacacacataactaaacctgaacg	att	tga
100740	96ORF008	2	8960..10201	413	tggcagaatttgggggataaacga	atg	tga
100741	96ORF009	2	17447..18670	407	gacgcaataacggaagtgcgtca	atg	tga
100742	96ORF010	1	38647..39819	390	taaatataaataaaggaggtgtgtaa	atg	tga
100743	96ORF011	-1	119..1195	358	gtagctcgctacccttattatttt	ttg	tga
100744	96ORF012	2	20045..21013	322	tttaatgacaaattacctgacatag	atg	tga
100745	96ORF013	3	29157..30098	313	acttattataaggaggtttgttag	ttg	taa
100746	96ORF014	1	21925..22839	304	agaaaaataaagtgaaggtataaaaa	atg	tag
100747	96ORF015	1	5812..6591	259	atacacggttaaagggtgggagaaatag	atg	taa
100748	96ORF016	1	7852..8607	251	aataaaatgttgaaaggagagaaaa	atg	taa
100749	96ORF017	3	3444..4190	248	aaatttaacattaataatcactttaa	gtg	taa
100750	96ORF018	-3	28281..29000	239	taagctatgttggaacatcgctagtc	atg	tga
100751	96ORF019	3	7188..7859	223	tttaccgttctaggacgtggtttaa	atg	taa
100752	96ORF020	3	21324..21908	194	gaagggcaaaaaggaggtttgatgat	atg	taa
100753	96ORF021	3	6612..7175	187	attaaaaatattaaaaggacggt	ata	tag
100754	96ORF022	2	24536..25093	185	aaagaaaaacgaaggagtgatttaa	atg	taa
100755	96ORF023	1	5275..5811	178	catgaaatggtaggaggtatgaaaa	gtg	tag
100756	96ORF024	3	14481..15014	177	taaaacgataggagataacgaataa	atg	taa
100757	96ORF025	2	25157..25666	169	ataaaaaaattgaaaggaggtatat	att	taa
100758	96ORF026	-3	15084..15590	168	tcattcttaacatagcccttaattc	atg	tga
100759	96ORF027	-1	1229..1732	167	aatagcaataaaggagtgtaaaac	atg	taa
100760	96ORF028	1	16960..17454	164	aaggcggtgtgatacagtgaaacaa	ttg	taa
100761	96ORF029	-1	1736..2227	163	tatgagaaaaaggagtcataataaaag	atg	taa
100762	96ORF030	1	25531..25995	154	ttttcaaggaggagagtcgctcgta	ctg	tag
100763	96ORF031	2	23633..24097	154	tttagtattgaaaggtgattctgtag	atc	tag
100764	96ORF032	-2	2248..2706	152	ataagacaccaaaagggtttggcgc	atg	tga
100765	96ORF033	-3	39147..39605	152	agcatataaatcggttagtggtttgt	ttg	taa
100766	96ORF034	2	13181..13615	144	tagaagtcgaaaaagtgaggcaat	ata	taa
100767	96ORF035	2	10628..11053	141	gagctaggattgcaagcaacgatat	ttg	tga
100768	96ORF036	2	24110..24535	141	gtatttttcatagaggtggttaaat	atg	taa
100769	96ORF037	1	12583..12996	137	atgaggaacagaaagcaaccaacttt	att	tga
100770	96ORF038	1	15628..16032	134	atgttaagaatgatgcctagttaa	ttg	taa
100771	96ORF039	3	39816..40220	134	ctaatacactttacttaataaaggg	gtg	taa
100772	96ORF040	-3	27528..27932	134	tttccataaataaacgaggacacca	atg	tga
100773	96ORF041	3	16206..16607	133	gatgagggcggaggtgtcagagtag	atg	tga
100774	96ORF042	2	35720..36106	128	aagttactataaactaaaattatggg	gtg	taa
100775	96ORF043	-2	35713..36081	122	ttaaacgtccccctcagttattgtt	ttg	taa
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100777	96ORF045	-3	5139..5504	121	ttctttttgtattctgtaatttca	att	tga
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100782	96ORF050	-2	2728..3072	114	tggtaaattagtattacattaagta	ata	taa
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100784	96ORF052	-1	20882..21211	109	gtagcaaaagagacaactaaaaaagt	gtg	taa
100785	96ORF053	1	40252..40578	108	acgactaattttttagtcggttttt	att	tag
100786	96ORF054	1	4942..5262	106	aatataaaactaaaaaaacaaaattt	atg	tag
100787	96ORF055	-2	4840..5151	103	ccgtcgcaatatatagttcgcttaa	atc	taa
100788	96ORF056	3	36324..36623	99	aatttaacacaaagttagtgccgta	atg	taa
100789	96ORF057	2	1394..1690	98	cttcagtggtcttttagcatttaa	ata	taa
100790	96ORF058	-3	26247..26537	96	tacttcttttctcataatctgacca	att	tga
100791	96ORF059	-1	21485..21772	95	agactcaacgcctttttgaaacatac	ttg	tga
100792	96ORF060	-3	22647..22931	94	cctcttttgtaaccgacaagactgta	ata	taa
100793	96ORF061	1	14023..14304	93	ttatctaattaaagggggacagtgta	gtg	taa
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100829	96ORF097	-1	36047..36265	72	tcaagcattacacctgtgacttttc	atc	taa
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101039	96ORF307	-1	20147..20263	38	ttgtacattatagcagtttaactcctt	att	tag
101040	96ORF308	-2	38158..38274	38	ttccgtatccactttctaaagaaagc	gtg	tga
101041	96ORF309	-2	35149..35265	38	agcttggtttgtatcgcttttaacga	ata	taa
101042	96ORF310	-2	31423..31539	38	gtaatatgattagggtctcctcttat	ttg	taa
101043	96ORF311	-2	10438..10554	38	cgcttttaaatcggttttaggtcact	atc	taa
101044	96ORF312	-2	1390..1506	38	gagaaacaacacaaacattaacaaca	atc	taa
101045	96ORF313	-3	33051..33167	38	acgtcctgtttcttagatcgtaatac	ata	tag
101046	96ORF314	-3	25194..25310	38	agcaaacggcttaagataaacttga	atc	taa
101047	96ORF315	-3	6273..6389	38	cattcttgcttaacacgtcagattga	ctg	tga
101048	96ORF316	-3	4281..4397	38	ataattcgatttcattaatcattaa	att	tag
101049	96ORF317	1	2260..2373	37	atgactcctttctcatatttcttt	ata	taa
101050	96ORF318	2	21230..21343	37	atttcacacttttttagttgtctct	ttg	taa
101051	96ORF319	3	18018..18131	37	atactgagtcaccaatttaagctcg	atg	tag
101052	96ORF320	3	36972..37085	37	attacagatatccctaagggtttccg	att	taa
101053	96ORF321	-1	36302..36415	37	ctcttgagttttttgacctaattta	atc	taa
101054	96ORF322	-1	32606..32719	37	ccataagttatttctccagttctat	att	taa
101055	96ORF323	-1	11453..11566	37	ttaaacgggtcttttttatcaattc	att	tga
101056	96ORF324	-1	7268..7381	37	tactgggtcgcgccagtggaagtctt	ata	tga
101057	96ORF325	-2	32347..32460	37	ttactgcattttgtatatggcgataa	atc	tag
101058	96ORF326	-2	24682..24795	37	acgtttattacgctcataaagccat	ata	tag
101059	96ORF327	-2	23905..24018	37	aaatggctgtggcgcttgaccatat	gtg	taa
101060	96ORF328	-2	21460..21573	37	agagcactaatacgtttttgttctt	ctg	tga
101061	96ORF329	-2	21208..21321	37	gacttaacttcttcgatattcattat	atc	tga
101062	96ORF330	-2	18085..18198	37	ccagtcgacaccagcaagttattct	ttg	tag
101063	96ORF331	-2	8170..8283	37	actttgagacgtcgtctgtctctct	atg	tag
101064	96ORF332	-2	5971..6084	37	caatttggttttcggttttctcttag	ttg	tag
101065	96ORF333	-3	37632..37745	37	accttgcttaatacaagtctgaattta	att	tga
101066	96ORF334	-3	29628..29741	37	ctgagttagtggttgaataatgtcat	ttg	tag
101067	96ORF335	-3	7164..7277	37	ttagcggatataccggttttctagtaa	atc	taa
101068	96ORF336	1	22903..23013	36	gtaaaaaaagacaatatgactatta	ctg	tga
101069	96ORF337	1	43258..43368	36	taattgacgtgggttatttttttaggt	ttg	taa
101070	96ORF338	2	12668..12778	36	gaactgggtggaatgggcattggaaca	atc	tag
101071	96ORF339	2	28292..28402	36	ttcactgctttaattcagttgtctta	ctg	taa
101072	96ORF340	2	35396..35506	36	ttcctaatagaacataagtcaacggg	att	tga
101073	96ORF341	3	25428..25538	36	actcgagaacaattagaaaaagcaa	ttg	tga
101074	96ORF342	-1	40913..41023	36	tatctgggaaatttaattctaataaaa	ata	tga
101075	96ORF343	-1	39173..39283	36	tgccacatttttagtgtaggattga	ttg	taa
101076	96ORF344	-1	37580..37690	36	gggtctacctttaacgtcgttttcag	ata	taa
101077	96ORF345	-1	31556..31666	36	ggattattctttctaataacttcaa	ttg	tga
101078	96ORF346	-1	29972..30082	36	ggctactccttattctaaaataataat	ttg	taa
101079	96ORF347	-1	28787..28897	36	ctgccaaagtctgtagcaattactt	ttg	tga
101080	96ORF348	-1	21839..21949	36	ttaaaaatccgataaaaataacattgc	ctg	tga
101081	96ORF349	-1	3647..3757	36	taaaacttccgaagttacccagcgt	ttg	tga

101082	96ORF350	-2	40801..40911	36	accattccaattttgcccataatgat	gtg	tag
101083	96ORF351	-2	38953..39063	36	tatctttttaaattctcgtaatagc	atc	taa
101084	96ORF352	-2	31585..31695	36	tagctgtcatcactagtatttttga	atc	taa
101085	96ORF353	-2	24550..24660	36	atagtcctgttttaccgcctcgtact	att	tag
101086	96ORF354	-2	20083..20193	36	atcatcatttttgatatttctcaaac	ata	tga
101087	96ORF355	-2	991..1101	36	gcattcttggcagtagcagcgtaaaac	atc	tag
101088	96ORF356	-3	38148..38258	36	taagaaagcgtgcgcgatcaataa	att	tga
101089	96ORF357	-3	8790..8900	36	tgaagtattctagcgtatttttct	ttg	tag
101090	96ORF358	-3	4458..4568	36	ttcataaaagtattctttgtagtat	atg	tag
101091	96ORF359	1	4666..4773	35	ttatcaaaatatacaacttaattaa	atc	tag
101092	96ORF360	1	11569..11676	35	ataaatttaccgaacatgaaaatga	att	tga
101093	96ORF361	2	6122..6229	35	ggaaaaacaaattgatgtgttagtga	ttg	taa
101094	96ORF362	-1	40418..40525	35	ttcgtagggtgtcattacttctttaa	ttg	tag
101095	96ORF363	-1	34358..34465	35	gttttgccttgatttcgatttgttga	atg	tga
101096	96ORF364	-1	20654..20761	35	ctatttccactgattccccatctaa	atg	tga
101097	96ORF365	-1	8423..8530	35	tcttttttagagttacgaggtttca	att	tag
101098	96ORF366	-1	2402..2509	35	tgacgtatggcaacatttttagatca	atc	taa
101099	96ORF367	-2	36607..36714	35	aaaataaaaagccagtgccgaagca	ctg	tag
101100	96ORF368	-2	27061..27168	35	caaatcgctcctgcagcgttcaataa	atc	tag
101101	96ORF369	-2	26470..26577	35	atgagttgttaagtttaccocaaat	atc	taa
101102	96ORF370	-2	10327..10434	35	ccgtgccattctctcggtataagta	ata	taa
101103	96ORF371	-2	8650..8757	35	gggtacgggttggttactgttgatat	atc	taa
101104	96ORF372	-3	14382..14489	35	gttcttttaattgatctactgttaa	att	taa
101105	96ORF373	-3	8151..8258	35	atgtttgttagtctctgtgtagtct	atg	taa
101106	96ORF374	-3	5007..5114	35	aaacgatttaagtggaaacattatc	ata	taa
101107	96ORF375	2	30563..30667	34	cgattagaaatctttaaanaagac	ttg	tga
101108	96ORF376	-1	19916..20020	34	tctatgtcaggttaatttgcattaa	att	taa
101109	96ORF377	-1	9236..9340	34	cttttctgttagtaattgtttttaa	atc	taa
101110	96ORF378	-1	9026..9130	34	actctttatctttagttgcttttaa	ata	tag
101111	96ORF379	-2	28447..28551	34	cttttgtgataataaagtttagtgc	ttg	tga
101112	96ORF380	-3	40329..40433	34	ccatttaccttcttgagatgttgga	ttg	tga
101113	96ORF381	-3	39801..39905	34	caaaagatgaaggcttttccatac	ttg	taa
101114	96ORF382	-3	33831..33935	34	atgttgttttgaactcgattaaagt	atc	tga
101115	96ORF383	-3	33687..33791	34	gttattacgtcttaatacttgtgtt	gtg	tag
101116	96ORF384	-3	13530..13634	34	tatacgcactagtactgatcactga	ttg	taa
101117	96ORF385	-3	3843..3947	34	tttgattgattgttctagtttaagaa	att	taa
101118	96ORF386	1	12256..12357	33	agtcataaagaagttagcaatgtga	ttg	tag
101119	96ORF387	2	2207..2308	33	tccaagactctttaactgttaactt	atc	tag
101120	96ORF388	2	2519..2620	33	attgttgaatttcgattgatctaaa	atg	tga
101121	96ORF389	2	22517..22618	33	agaagtaaaatgcgtaatgctttag	atg	tag
101122	96ORF390	2	27302..27403	33	ttccaaaattgggctaatagtgtgag	ctg	taa
101123	96ORF391	2	32384..32485	33	actaaaaaggttgagaaagctgtga	atg	taa
101124	96ORF392	2	39287..39388	33	aaaaacggtagctgtagtatcaatca	atc	tag
101125	96ORF393	3	18153..18254	33	gtagtatatgccgactttgatttga	atg	taa
101126	96ORF394	3	24189..24290	33	tcagaccctaaccattacaaaactag	ttg	tga
101127	96ORF395	-1	15266..15367	33	tcgataatttgtatagcttgtttta	atg	tag
101128	96ORF396	-2	32239..32340	33	tttttagtgaaagcatctagtgttga	ata	tag
101129	96ORF397	-2	16123..16224	33	ttatgtgtgcctatcatattacaa	ttg	tag
101130	96ORF398	-2	13648..13749	33	tctttaactgaatgttgaatagcat	ttg	tag
101131	96ORF399	-2	10987..11088	33	acttctgtaggattcttatatcaa	ttg	tga
101132	96ORF400	-2	3382..3483	33	cttactggtaattcttcaaaattaa	atg	taa
101133	96ORF401	-3	40794..40895	33	ccatattgatgtgaaagtgtttaaat	ttg	taa
101134	96ORF402	-3	39978..40079	33	atattcctaataactgaacctaata	att	tga
101135	96ORF403	-3	38607..38708	33	atcttcagtgtaaaatcgacagcca	atg	tag
101136	96ORF404	-3	21288..21389	33	cagacaccgtcttaagtccctttag	ata	taa

Table 11

SEQUENCE INFORMATION FOR PHAGES MATCHING WITH TABLE 1

M32695

Bacteriophage PM2 nuclease cleavage site

gi|166145|gb|M32695|BM2NCS [166145]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M32693

Bacteriophage PM2 Hind III fragment 4

gi|166144|gb|M32693|BM24HIND3 [166144]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M32693

Bacteriophage PM2 Hind III fragment 4

gi|166144|gb|M32693|BM24HIND3 [166144]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M32694

Bacteriophage PM2 Hind III fragment 3

gi|166143|gb|M32694|BM23HIND3 [166143]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M26134

Bacteriophage PM2 structural protein gene containing purine/pyrimidine rich regions and anti-Z-DNA-IgG binding regions, complete cds

gi|289360|gb|M26134|BM2PROTIV [289360]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

J02452

bacteriophage fi 3'-terminal region ma

gi|215409|gb|J02452|PFITR3 [215409]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

AF020798

Bacteriophage Chp1 genome DNA, complete sequence

gi|217761|dbj|D00624|BCP1 [217761]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 12 protein links, or 1 genome link)

X72793

Clostridium botulinum C phage BONT/C1, ANTP-139, ANTP-33, ANTP-17, ANTP-70 genes and ORF-22

gi|516171|emb|X72793|CBCBONT [516171]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 6 protein links, or 4 nucleotide neighbors)

X51464

Clostridium botulinum D Phage C3 gene for exoenzyme C3

gi|14907|emb|X51464|CBDPE3 [14907]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

D90210

Bacteriophage c-st (from C. botulinum) C1-tox gene for botulinum C1 neurotoxin

gi|217780|dbj|D90210|CSTC1TOX [217780]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

S49407

type D neurotoxin [bacteriophage d-16 phi, host = C. botulinum, type D, CB16, Genomic, 4087 nt]
gi|260238|gb|S49407|S49407 [260238]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

X53370

Bacteriophage phi29 temperature sensitive mutant TS2(98) DNA polymerase gene
gi|15733|emb|X53370|POTS298 [15733]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 7 nucleotide neighbors)

X53371

Bacteriophage phi29 temperature sensitive mutant TS2(24) DNA polymerase gene
gi|15731|emb|X53371|POTS224 [15731]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 7 nucleotide neighbors)

X05973

Bacteriophage phi29 prohead RNA
gi|15680|emb|X05973|POP29PRO [15680]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 4 nucleotide neighbors)

V01155

Left end of bacteriophage phi-29 coding for 15 potential proteins Among these are the terminal protein and the proteins encoded by the genes 1, 2 (sus), 3, and (probably) 4
gi|15659|emb|V01155|POP29B [15659]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 16 protein links, or 16 nucleotide neighbors)

X73097

Bacteriophage phi-29 left origin of replication
gi|312194|emb|X73097|BP29ORIL [312194]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 5 nucleotide neighbors)

M14430

Bacteriophage phi-29 gene-17 gene, complete cds
gi|215321|gb|M14430|P29G17A [215321]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 6 protein links, or 8 nucleotide neighbors)

M14431

Bacteriophage phi-29 gene-16 gene, complete cds
gi|215319|gb|M14431|P29G16A [215319]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 7 nucleotide neighbors)

M20693

Bacteriophage phi-29 DNA, 3' end
gi|215343|gb|M20693|P29REPINB [215343]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 4 nucleotide neighbors)

M21016

Bacteriophage phi-29 DNA, 5' end
gi|215342|gb|M21016|P29REPINA [215342]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M12456

Bacteriophage phi-29 genes 9, 10 and 11 encoding p9 tail, incomplete, p10 connector, complete, and p11 lower collar, incomplete, respectively

gi|215338|gb|M12456|P29P9 [215338]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

M14782

Bacillus phage phi-29 head morphogenesis, major head protein, head fiber protein, tail protein, upper collar protein, lower collar protein, pre-neck-appendage protein, morphogenesis(13), lysis, morphogenesis(15), encapsidation genes, complete cds
gi|215323|gb|M14782|P29LATE2 [215323]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 11 protein links, or 11 nucleotide neighbors)

M26968

Bacteriophage phi-29 (from Bacillus subtilis) proteins p1 delta-1 genes, complete cds, and the sus1(629) mutation

gi|341558|gb|M26968|P29P1D1A [341558]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

J02448

Bacteriophage f1, complete genome

gi|166201|gb|J02448|F1CCG [166201]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 10 protein links, 205 nucleotide neighbors, or 1 genome link)

M24832

Bacteriophage f2 coat protein gene, partial cds

gi|166228|gb|M24832|F2CRNACA [166228]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

J02451

Bacteriophage fd, strain 478, complete genome

gi|215394|gb|J02451|PFDCG [215394]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,5 MEDLINE links, 10 protein links, 204 nucleotide neighbors, or 1 genome link)

M34834

Bacteriophage fr replicase gene, 5' end

gi|166139|gb|M34834|BFRREGRA [166139]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 9 nucleotide neighbors)

M38325

Bacteriophage fr replicase gene, 5' end

gi|166137|gb|M38325|BFRREGR [166137]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 9 nucleotide neighbors)

M35063

Bacteriophage fr coat protein replicase cistron (R region) RNA

gi|166134|gb|M35063|BFRRCRRA [166134]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 3 nucleotide neighbors)

S66567

alpha-atrial natriuretic factor/coat protein=fusion polypeptide [human, bacteriophage fr, expression vector pFAN15, PlasmidSyntheticRecombinant, 510 nt]

gi|435742|gb|S66567|S66567 [435742]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 15 nucleotide neighbors)

X15031

Bacteriophage fr RNA genome

gi|15071|emb|X15031|LEBFRX [15071]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, 9 nucleotide neighbors, or 1 genome link)

U51233

Mus musculus neutralizing anti-RNA-bacteriophage fr immunoglobulin variable region light chain (IgM) mRNA, partial cds

gi|1277150|gb|U51233|MMU51233 [1277150]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 1669 nucleotide neighbors)

U51232

Mus musculus neutralizing anti-RNA-bacteriophage fr immunoglobulin variable region heavy chain (IgM) mRNA, partial cds

gi|1277148|gb|U51232|MMU51232 [1277148]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 1073 nucleotide neighbors)

U02303

Bacteriophage If1, complete genome

gi|3676280|gb|U02303|B2U02303 [3676280]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 10 protein links, or 1 genome link)

V00604

Phage M13 genome

gi|14959|emb|V00604|INM13X [14959]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, or 205 nucleotide neighbors)

A32252

Synthetic bacteriophage M13 protein III probe

gi|1567340|emb|A32252|A32252 [1567340]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

A32251

Synthetic bacteriophage M13 protein III probe

gi|1567339|emb|A32251|A32251 [1567339]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

M12465

Bacteriophage M13 mp10 mutations in lac operon

gi|215210|gb|M12465|M13LACMUT [215210]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 215 nucleotide neighbors)

M24177

Synthetic Bacteriophage M13 (clone M13.SV.B12) SV40 early promoter region DNA

gi|209416|gb|M24177|SYNSVB12 [209416]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)

M24176

Synthetic Bacteriophage M13 (clone M13.SV.B11) SV40 early promoter region DNA

gi|209415|gb|M24176|SYNSVB11 [209415]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)

M24175

Synthetic Bacteriophage M13 (clone M13.SV.8) SV40 early promoter region DNA
gi|208806|gb|M24175|SYNM13SV8 [208806]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 242 nucleotide neighbors)

M19979

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33
gi|207813|gb|M19979|SYN33M13M [207813]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 617 nucleotide neighbors)

M19565

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33
gi|207808|gb|M19565|SYN33M13H [207808]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 567 nucleotide neighbors)

M19564

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33
gi|207807|gb|M19564|SYN33M13G [207807]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 12 nucleotide neighbors)

M19563

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33
gi|207806|gb|M19563|SYN33M13F [207806]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 262 nucleotide neighbors)

M19561

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33
gi|207804|gb|M19561|SYN33M13D [207804]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 27 nucleotide neighbors)

M19560

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33
gi|207803|gb|M19560|SYN33M13C [207803]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M19559

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33
gi|207802|gb|M19559|SYN33M13B [207802]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 227 nucleotide neighbors)

M10568

Bacteriophage M13 replicative form II, replication origin, specific nick location
gi|215220|gb|M10568|M13ORIB [215220]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 650 nucleotide neighbors)

M10910

Bacteriophage M13 gene II regulatory region and M13sj1 mutant
gi|215209|gb|M10910|M13IIREG [215209]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 72 nucleotide neighbors)

M38295

Bacteriophage M13 HaeIII restriction fragment DNA
gi|215208|gb|M38295|M13HAEIII [215208]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 67 nucleotide neighbors)

E02067

DNA encoding a part of Bacteriophage M13 tg 127
 gi|2170311|dbj|E02067|E02067 [2170311]
 (View GenBank report, FASTA report, ASN.1 report, or Graphical view)

J02467

Bacteriophage MS2, complete genome
 gi|215232|gb|J02467|MS2CG [215232]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 8 MEDLINE links, 4 protein links, 20 nucleotide neighbors, or 1 genome link)

AJ004950

Bacteriophage P1 ban gene
 gi|3688226|emb|AJ011592|BP1011592 [3688226]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)

U88974

Bacteriophage P1 structural lytic transglycosylase (orf47), pep44b (orf44b),
 pep44a (orf44a), and pep43 (orf43) genes, complete cds; and pep42 (orf42) gene, partial cds
 gi|2661099|gb|AF035607|AF035607 [2661099]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 protein links, or 1 nucleotide neighbor)

AJ000741

Bacteriophage P1 darA operon
 gi|2462938|emb|AJ000741|BPAJ7641 [2462938]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, or 31 nucleotide neighbors)

X01828

Bacteriophage P1 recombinase gene cin
 gi|15133|emb|X01828|MYP1CIN [15133]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

X98146

Bacteriophage P1 DNA sequence around the Op88 operator
 gi|1359513|emb|X98146|BP1OP88OP [1359513]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor)

S61175

immI operon: icd=cell division repressor, ant1=antirepressor (promoters
 P51a, P51b) [bacteriophage P1, Genomic, 728 nt]
 gi|385908|gb|S61175|S61175 [385908]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)

X87824

Bacteriophage P1 gene 26
 gi|861164|emb|X87824|XXBP1G26 [861164]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)

X15638

Phage P1 DNA for lytic replicon containing promoter P53 and two open reading frames
 gi|15735|emb|X15638|PP1LREP [15735]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 24 nucleotide neighbors)

X17512

Bacteriophage P1 DNA for immunity region immI

gi|15479|emb|X17512|P1IMMUNITY [15479]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 4 nucleotide neighbors)

X16005

Bacteriophage P1 cI gene for P1cI repressor protein

gi|15477|emb|X16005|P1C1 [15477]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

X03453

Bacteriophage P1 cre gene for recombinase protein

gi|15135|emb|X03453|MYP1CRE [15135]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors)

X06561

Bacteriophage P1 cI gene 5'-region

gi|15128|emb|X06561|MYP1C1 [15128]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 6 nucleotide neighbors)

V01534

Bacteriophage P1 genome fragment (IS2 insertion spot). This regions contains

four unidentified reading frames and is known as insertion hot spot for IS2 insertion sequences

gi|15118|emb|V01534|MYOVP1 [15118]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 3 nucleotide neighbors)

X56951

Bacteriophage P1 gene10

gi|406728|emb|X56951|BPP1GP10 [406728]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 3 protein links, or 1 nucleotide neighbor)

K02380

Bacteriophage P1 replication region including repA, parA, and parB genes and

incA, incB, and incC incompatibility determinants

gi|215652|gb|K02380|PP1REP [215652]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,5 MEDLINE links, 4 protein links, or 8 nucleotide neighbors)

X87674

Bacteriophage P1 lydA & lydB genes

gi|974763|emb|X87674|BACP1LYD [974763]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

X87673

Bacteriophage P1 gene 17

gi|974761|emb|X87673|BACP117 [974761]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

M16618

Bacteriophage P1 cI repressor binding sites

gi|215600|gb|M16618|PP1C1 [215600]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

SEG_PP1CTN

Bacteriophage P1 cin gene encoding recombinase, cixL recombination site, and 5' end of C invertible element
gi|215607|gb|SEG_PP1CTN [215607]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

K03173

Bacteriophage P1 C invertible element, right end, and cixR recombination site
gi|215606|gb|K03173|PP1CIN2 [215606]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

215605

Bacteriophage P1 cin gene encoding recombinase, cixL recombination site, and 5' end of C invertible element
gi|215605|lc|X01828 [215605]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M25470

Bacteriophage P1 tail fiber protein gene, complete cds
gi|341349|gb|M25470|PP1TFPR [341349]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)

M34382

Bacteriophage P1 sim region proteins, complete cds
gi|215661|gb|M34382|PP1SIM [215661]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 protein links)

M81956

Bacteriophage P1 R protein (R) gene, complete cds
gi|215658|gb|M81956|PP1RP [215658]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 4 nucleotide neighbors)

M37080

Bacteriophage P1 mini-P1 plasmid origin of replication
gi|215657|gb|M37080|PP1REPOR [215657]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 46 nucleotide neighbors)

M27041

Bacteriophage P1 ref gene, complete cds
gi|215650|gb|M27041|PP1REF [215650]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

L01408

Bacteriophage P1 partition protein (parB) gene, 3' end
gi|215642|gb|L01408|PP1PARB [215642]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 41 nucleotide neighbors)

SEG_PP1PAR

Bacteriophage miniplasmid P1 parA gene, 5' end
gi|215639|gb|SEG_PP1PAR [215639]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 48 nucleotide neighbors)

M36425

Bacteriophage miniplasmid P1 parB gene, 3' end
gi|215638|gb|M36425|PP1PAR2 [215638]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M36424

Bacteriophage miniplasmid P1 *parA* gene, 5' end
gi|215637|gb|M36424|PP1PAR1 [215637]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

M11129

Bacteriophage P1 miniplasmid origin of replication region
gi|215632|gb|M11129|PP1ORIM [215632]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 43 nucleotide neighbors)

M25414

Bacteriophage P1 *cI* repressor binding site, operator 88 (Op88)
gi|215631|gb|M25414|PP1OP88A [215631]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)

M25413

Bacteriophage P1 *cI* repressor binding site, operator 68 (Op68)
gi|215630|gb|M25413|PP1OP68A [215630]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

M25412

Bacteriophage P1 *cI* repressor binding site, operator 21 (Op21)
gi|215629|gb|M25412|PP1OP21A [215629]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)

M10510

Bacteriophage P1 recombination site *loxR*
gi|215628|gb|M10510|PP1LOXR [215628]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)

M10287

Bacteriophage P1 *loxP* X *loxP* recombination site
gi|215627|gb|M10287|PP1LOXPX [215627]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)

M10494

Bacteriophage P1 recombination site *loxP*
gi|215626|gb|M10494|PP1LOXP [215626]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 134 nucleotide neighbors)

M10511

Bacteriophage P1 recombination site *loxL*
gi|215625|gb|M10511|PP1LOXL [215625]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)

M10512

Bacteriophage P1 recombination site *loxB*
gi|215624|gb|M10512|PP1LOXB [215624]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

M10145

Bacteriophage P1 genome fragment with recombination site *loxP*
gi|215623|gb|M10145|PP1CREX [215623]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 21 nucleotide neighbors)

M13327

Bacteriophage P1 *Cin* recombinase activated cross over site, junction IV, clone pSHI326
gi|215622|gb|M13327|PP1CN26IV [215622]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)

M13325

Bacteriophage P1 *Cin* recombinase activated cross over site, junction II, clone pSHI326
gi|215621|gb|M13325|PP1CN26II [215621]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1401 nucleotide neighbors)

M13323

Bacteriophage P1 *Cin* recombinase activated cross over site, junction IV, clone pSHI325
gi|215620|gb|M13323|PP1CN25IV [215620]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)

M13321

Bacteriophage P1 *Cin* recombinase activated cross over site, junction II, clone pSHI325
gi|215619|gb|M13321|PP1CN25II [215619]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1058 nucleotide neighbors)

M13324

Bacteriophage P1 *Cin* recombinase activated cross over site, junction I, clone pSHI326
gi|215618|gb|M13324|PP1CIR26I [215618]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)

M13319

Bacteriophage P1 *Cin* recombinase activated cross over site, right junction, clone pSHI327
gi|215617|gb|M13319|PP1CIN27R [215617]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)

M13320

Bacteriophage P1 *Cin* recombinase activated cross over site, junction I, clone pSHI325
gi|215616|gb|M13320|PP1CIN25I [215616]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)

M13318

Bacteriophage P1 *Cin* recombinase activated cross over site, left junction, clone pSHI324
gi|215615|gb|M13318|PP1CIN24L [215615]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1370 nucleotide neighbors)

M13317

Bacteriophage P1 *Cin* recombinase activated cross over site, right junction, clone pSHI323
gi|215614|gb|M13317|PP1CIN23M [215614]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1055 nucleotide neighbors)

M13316

Bacteriophage P1 *Cin* recombinase activated cross over site, left junction, clone pSHI323
gi|215613|gb|M13316|PP1CIN23L [215613]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)

M13315

Bacteriophage P1 *Cin* recombinase activated cross over site, right junction, clone pSHI322
gi|215612|gb|M13315|PP1CIN22R [215612]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)

M13314

Bacteriophage P1 *Cin* recombinase activated cross over site, left junction, clone pSHI322

gi|215611|gb|M13314|PP1CIN22L [215611]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1401 nucleotide neighbors)

M13313

Bacteriophage P1 *Cin* recombinase activated cross over site, right junction, clone pSHI321

gi|215610|gb|M13313|PP1CIN21R [215610]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 7 nucleotide neighbors)

M13312

Bacteriophage P1 *Cin* recombinase activated cross over site, left junction, clone pSHI321

gi|215609|gb|M13312|PP1CIN21L [215609]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1058 nucleotide neighbors)

M16568

Bacteriophage P1 *c4* repressor gene, complete cds

gi|215603|gb|M16568|PP1C4 [215603]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

M13326

Bacteriophage P1 *Cin* recombinase activated cross over site, junction III, clone pSHI326

gi|215602|gb|M13326|PP1C26III [215602]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1192 nucleotide neighbors)

M13322

Bacteriophage P1 *Cin* recombinase activated cross over site, junction III, clone pSHI325

gi|215601|gb|M13322|PP1C25III [215601]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1231 nucleotide neighbors)

J05651

Bacteriophage P1 modulator protein (*bof*) gene, complete cds

gi|215598|gb|J05651|PP1BOFY1 [215598]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

M33224

Bacteriophage P1 regulatory protein (*bof*) gene, complete cds

gi|215596|gb|M33224|PP1BOFFO [215596]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

M10288

E.coli/bacteriophage P1 *loxR* recombination site

gi|146647|gb|M10288|ECOLOXR [146647]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 nucleotide neighbors)

M10289

E.coli/bacteriophage P1 *loxL* recombination site

gi|146646|gb|M10289|ECOLOXL [146646]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M10290

E.coli *loxB* site, which can recombine with bacteriophage P1 *loxP* site

gi|146645|gb|M10290|ECOLOXB [146645]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M10287

Bacteriophage P1 loxP X loxP recombination site

gi|215627|gb|M10287|PP1LOXPX [215627]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)

M74046

Bacteriophage P1 pacA and pacB genes, complete cds

gi|215634|gb|M74046|PP1PACAB [215634]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

M95666

Bacteriophage P1 gene 10, doc and phd genes, complete cds

gi|463276|gb|M95666|PP1PHDDOC [463276]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 4 protein links, or 1 nucleotide neighbor)

M25604

Bacteriophage Q-beta mutated autonomously replicating sequence MDV1 RNA fragment

gi|556359|gb|M25604|PQBARSMT [556359]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors)

V00643

first half of the phage Q-beta gene for coat protein

gi|15088|emb|V00643|LEQBET [15088]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

M25167

Bacteriophage Q-beta RNA fragment recovered from replicase binding complex

gi|556362|gb|M25167|PQBREPLICB [556362]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)

M24876

Bacteriophage Q-beta replicase RNA, 5' end

gi|556360|gb|M24876|PQBREPLICA [556360]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

M25444

Synthetic bacteriophage Q-beta DNA

gi|209118|gb|M25444|SYNPQBTERM [209118]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors)

M25463

Bacteriophage Q-beta self-replicating microvariant (+) RNA

gi|532489|gb|M25463|PQBMVSRRA [532489]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

M25014

Bacteriophage Q-beta RNA replicase gene, 5' end, and maturation protein gene, 3' end

gi|294316|gb|M25014|PQBREPLC [294316]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

M25065

Bacteriophage Q-beta RNA sequence with putative stem loop

gi|294315|gb|M25065|PQBLOOP [294315]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)

M10265

Bacteriophage Q-beta RNA molecule with the ability to replicate extracellularly
gi|215726|gb|M10265|PQBRNA [215726]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors)

M24815

Bacteriophage Q-beta specified replicase subunit RNA,
gi|215725|gb|M24815|PQBREPL [215725]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors)

M25461

Bacteriophage Q-beta plus-strand RNA, 5' terminus
gi|215724|gb|M25461|PQBPS5E [215724]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

M25462

Bacteriophage Q-beta plus-strand RNA, 3' terminus
gi|215723|gb|M25462|PQBPS3E [215723]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 8 nucleotide neighbors)

M24871

Bacteriophage Q-beta nanovariant WSIII RNA
gi|215722|gb|M24871|PQBNVWSIC [215722]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)

M24870

Bacteriophage Q-beta nanovariant WSII RNA
gi|215721|gb|M24870|PQBNVWSIB [215721]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)

M24869

Bacteriophage Q-beta nanovariant WSI RNA
gi|215720|gb|M24869|PQBNVWSIA [215720]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)

M10495

Coliphage Q-beta MDV-1(+) RNA
gi|215719|gb|M10495|PQBMDV1A [215719]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 10 nucleotide neighbors)

J02484

bacteriophage qbeta coat protein cistron first half
gi|215717|gb|J02484|PQBCP5 [215717]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

M57754

Bacteriophage Q-beta minus strand RNA, 5' terminus
gi|215716|gb|M57754|PQBBMS5E [215716]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 8 nucleotide neighbors)

M24297

Bacteriophage Q-beta 5'-terminal region of the minus strand
gi|215715|gb|M24297|PQB5END [215715]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 8 nucleotide neighbors)

M10695

218

Bacteriophage Q-beta, MDV-1 RNA

gi|215714|gb|M10695|PQB1R [215714]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 12 nucleotide neighbors)

M24827

Bacteriophage R17 A protein gene, 5' end

gi|216078|gb|M24827|R17RNACIS [216078]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 5 nucleotide neighbors)

M24829

Bacteriophage R17 coat protein gene, 5' end

gi|216075|gb|M24829|R17CP5 [216075]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 5 nucleotide neighbors)

J02488

bacteriophage r17 rna synthetase initiation site

gi|216080|gb|J02488|R17RNASYN [216080]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 2 protein links, or 6 nucleotide neighbors)

J02487

bacteriophage r17 coat protein initiation site

gi|216073|gb|J02487|R17COATP [216073]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

J02486

bacteriophage r17 a protein initiation site

gi|216071|gb|J02486|R17APROT [216071]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M24826

Bacteriophage R17 coat protein RNA fragment

gi|216077|gb|M24826|R17CPRAA [216077]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 7 nucleotide neighbors)

M24296

Bacteriophage R17 3'-terminal fragment A RNA

gi|216070|gb|M24296|R173TFA [216070]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 9 nucleotide neighbors)

1TFN

structure refinement for a 24-nucleotide rna hairpin, nmr, minimized average

structure ribonucleic acid, hairpin, bacteriophage r17 mol_id: 1; molecule: r17c; chain: null; engineered: yes

gi|1942336|pdb|1TFN| [1942336]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 structure link)

1RPEA

rna (5'-d(gpgpgpapcpupgpgpapcpupcpapcpup cpapgpupcpupapu-3') (24-mer rna

hairpin coat protein binding site for bacteriophage r17) (nmr, minimized average structure)

gi|1421020|pdb|1RHT| [1421020]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 structure link)

M14428

Bacteriophage S13 circular DNA, complete genome

gi|216089|gb|M14428|S13CG [216089]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 12 protein links, 26 nucleotide neighbors, or 1 genome link)

J05393

Bacteriophage T1 DNA N-6-adenine-methyltransferase (M.T1) gene, complete cds

gi|166163|gb|J05393|BT1NAMTA [166163]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

L46845

Bacteriophage T2 frd3, frd2 genes, complete cds

gi|951387|gb|L46845|PT2FRD32G [951387]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 17 nucleotide neighbors)

L43611

Bacteriophage T2 fibrin (wac) gene, complete cds

gi|903869|gb|L43611|PT2WAC [903869]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)

M24812

Bacteriophage T2 secondary structure RNA sequence

gi|215796|gb|M24812|PT2RNA [215796]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors)

M22342

Bacteriophage T2 DNA-(adenine-N6)methyltransferase (dam) gene, complete cds

gi|215792|gb|M22342|PT2DAM [215792]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

S57515

orf 61.2 {intergenic region between 41 and 61} [bacteriophage T2, Genomic, 323 nt]

gi|298524|gb|S57515|S57515 [298524]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

X05312

Bacteriophage T2 gene 38 for receptor recognizing protein

gi|15197|emb|X05312|MYT2G38 [15197]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

X04442

Bacteriophage T2 gene 37 for receptor recognizing protein

gi|15195|emb|X04442|MYT2G37 [15195]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

X12460

Bacteriophage T2 gene 32 mRNA for single-stranded DNA binding protein

gi|15192|emb|X12460|MYT2G32 [15192]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 14 nucleotide neighbors)

X57797

Bacteriophage T2 gene for gp12

gi|14875|emb|X56555|BT2GP12 [14875]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 2 nucleotide neighbors)

X01755

Bacteriophage T2 tail fiber gene 36

gi|15189|emb|X01755|MYT2F36 [15189]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

M14784

Bacteriophage T3 strain amNG220B right end, tail fiber protein, lysis protein and DNA packaging proteins, complete cds

gi|215810|gb|M14784|PT3RE [215810]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 9 protein links, or 10 nucleotide neighbors)

SEG_PT3RNAPOL

Bacteriophage T3 RNA polymerase III gene, 5' end

gi|710559|gb|SEG_PT3RNAPOL [710559]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

M22610

Bacteriophage T3 RNA polymerase III gene, 3' end

gi|340722|gb|M22610|PT3RNAPOL2 [340722]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M22609

Bacteriophage T3 RNA polymerase III gene, 5' end

gi|340721|gb|M22609|PT3RNAPOL1 [340721]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

X05031

Bacteriophage T3 gene region 1-2.5 with primary origin of replication

gi|15719|emb|X05031|POT3ORI [15719]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 11 protein links, or 5 nucleotide neighbors)

X03964

Bacteriophage T3 early control region pos. 308-810 from genome left end

gi|15718|emb|X03964|POT3EP [15718]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 20 nucleotide neighbors)

X17255

Bacteriophage T3 gene 1 to gene 11

gi|15682|emb|X17255|POT3111G [15682]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,4 MEDLINE links, 36 protein links, 17 nucleotide neighbors, or 1 genome link)

X15840

Phage T3 gene 10

gi|15625|emb|X15840|PODT3G10 [15625]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 nucleotide neighbors)

X02981

Bacteriophage T3 gene 1 for RNA polymerase

gi|15561|emb|X02981|PODOT3P [15561]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

J02503

bacteriophage t3 5' end, terminally redundant sequence (trs)

gi|215816|gb|J02503|PT3TRS1 [215816]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

SEG_PT3TRS

bacteriophage λ 3' end, terminally redundant sequence (trs)

gi|215818|gb|SEG_PT3TRS [215818]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

J02504

bacteriophage λ 3' end, terminally redundant sequence (trs)

gi|215817|gb|J02504|PT3TRS2 [215817]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

H YPERLINK <http://www.rs.noda.sut.ac.jp/~kunisawa> h t t p://www.rs.noda.sut.ac.jp/~kunisawa
Bacteriophage T4 genomic database compiled by Arisaka et al.

X95646

Bacteriophage T5 DNA for region 60.5%-71% of the T5 genome

gi|2791557|emb|AJ001191|BTJ001191 [2791557]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 7 MEDLINE links, 12 protein links, or 6 nucleotide neighbors)

X56847

Bacteriophage T5 genomic region encoding early genes D10-D15

gi|15407|emb|X12930|MYT5D10 [15407]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 5 protein links, or 4 nucleotide neighbors)

AF039886

Bacteriophage T5 subclone T5.5.3r5.18r, single pass sequence, genomic survey sequence

gi|2811154|gb|AF039886|AF039886 [2811154]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF039885

Bacteriophage T5 subclone T5.40f,41f, single pass sequence, genomic survey sequence

gi|2811153|gb|AF039885|AF039885 [2811153]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF039884

Bacteriophage T5 subclone T5.26.fr, single pass sequence, genomic survey sequence

gi|2811152|gb|AF039884|AF039884 [2811152]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF039883

Bacteriophage T5 subclone 10-T5.5.7F, single pass sequence, genomic survey sequence

gi|2811151|gb|AF039883|AF039883 [2811151]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF039882

Bacteriophage T5 subclone 41-T5.5.4BF, single pass sequence, genomic survey sequence

gi|2811150|gb|AF039882|AF039882 [2811150]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF039881

Bacteriophage T5 subclone 39-T5.5.4aF, single pass sequence, genomic survey sequence

gi|2811149|gb|AF039881|AF039881 [2811149]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor)

AF039880

Bacteriophage T5 subclone 19-T5.7.2r, single pass sequence, genomic survey sequence
gi|2811148|gb|AF039880|AF039880 [2811148]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039879

Bacteriophage T5 subclone 18-T5.7.2F, single pass sequence, genomic survey sequence
gi|2811147|gb|AF039879|AF039879 [2811147]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039878

Bacteriophage T5 subclone 11-T5.5.7R, single pass sequence, genomic survey sequence
gi|2811146|gb|AF039878|AF039878 [2811146]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 nucleotide neighbors)

AF039877

Bacteriophage T5 subclone T5.4FR, single pass sequence, genomic survey sequence
gi|2811145|gb|AF039877|AF039877 [2811145]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039876

Bacteriophage T5 subclone 22-T5.16R, single pass sequence, genomic survey sequence
gi|2811144|gb|AF039876|AF039876 [2811144]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039875

Bacteriophage T5 subclone 21-T5.16R, single pass sequence, genomic survey sequence
gi|2811143|gb|AF039875|AF039875 [2811143]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039874

Bacteriophage T5 subclone 21-T5.16F, single pass sequence, genomic survey sequence
gi|2811142|gb|AF039874|AF039874 [2811142]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039873

Bacteriophage T5 subclone 09-T5.6F, single pass sequence, genomic survey sequence
gi|2811141|gb|AF039873|AF039873 [2811141]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039872

Bacteriophage T5 subclone 09-T5.6R, single pass sequence, genomic survey sequence
gi|2811140|gb|AF039872|AF039872 [2811140]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 nucleotide neighbors)

AF039871

Bacteriophage T5 subclone 04-T5.26.R, single pass sequence, genomic survey sequence
gi|2811139|gb|AF039871|AF039871 [2811139]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039870

Bacteriophage T5 subclone 13-T5.42F, single pass sequence, genomic survey sequence
gi|2811138|gb|AF039870|AF039870 [2811138]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

X69460

Bacteriophage T5 ltf gene for L-shaped tail fibers

gi|15415|emb|X69460|MYT5LTF [15415]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 1 protein link, or 4 nucleotide neighbors)

X03402

Bacteriophage T5 D15 gene for 5' exonuclease

gi|15413|emb|X03402|MYT5EXOG [15413]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

Z11972

Bacteriophage T5 tRNA-Tyr, tRNA-Glu, tRNA-Trp, tRNA-Phe, tRNA-Cys and tRNA-Asn genes, and ORFs 91aa, 90aa, 42aa and 172aa

gi|15795|emb|Z11972|T56TRNAG [15795]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 3 nucleotide neighbors)

X03898

Bacteriophage T5 genes for tRNA-His, -Ser and -Leu

gi|15786|emb|X03898|STT5RN1 [15786]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 MEDLINE links)

X04177

Bacteriophage T5 gene for transfer RNA-Gln

gi|15421|emb|X04177|MYT5TRNQ [15421]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

X03899

Bacteriophage T5 genes for tRNA-Val, -Lys, -fMet, -Pro and -Ile3

gi|15787|emb|X03899|STT5RN2 [15787]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

X03798

Bacteriophage T5 gene for tRNA-Asp (GUC)

gi|15472|emb|X03798|NCT5TRDG [15472]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

Y00364

Bacteriophage T5 tRNA gene cluster (27.8%-22.4%)

gi|15420|emb|Y00364|MYT5TRN [15420]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 13 nucleotide neighbors)

X03140

Bacteriophage T5 DNA with rho-dependent transcription terminator (Hind III-P fragment)

gi|15417|emb|X03140|MYT5RHO [15417]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

Z35070

Bacteriophage T6 DNA

gi|535228|emb|Z35074|MYEREGBT6 [535228]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

AF060870

Coliphage T6 small subunit distal tail fiber (gene 36) gene, partial cds; and large subunit distal tail fiber (gene 37) and tail fiber adhesin (gene 38) genes, complete cds

gi|3676458|gb|AF052605|AF052605 [3676458]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 protein links, or 2 nucleotide neighbors)

Z35072

Bacteriophage T6 DNA encoding ORF19.1 gene and g19 gene

gi|535232|emb|Z35072|MYTAILT6 [535232]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 protein links)

X12488

Bacteriophage T6 gene 32 mRNA for single-stranded DNA binding protein

gi|15843|emb|X12488|MYT6G32 [15843]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 14 nucleotide neighbors)

Z78095

Bacteriophage T6 DNA (1506 bp)

gi|1488562|emb|Z78095|BPHZ78095 [1488562]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 4 nucleotide neighbors)

Z35079

Bacteriophage T6 DNA for Ip5, Ip6

gi|535215|emb|Z35079|MY57BT6 [535215]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

X68725

E.coli bacteriophage T6 gene for beta-glucosyl-HMC-alpha-glucosyl-transferase

gi|296439|emb|X68725|ECT6 [296439]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

X69894

Bacteriophage T6 alt gene for ADP-Ribosyltransferase

gi|15422|emb|X69894|MYT6ADP [15422]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

L46846

Bacteriophage T6 frd3, frd2 genes, complete cds

gi|951390|gb|L46846|PT6FRD32G [951390]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 protein links)

M27738

Bacteriophage T6 translational repressor protein (regA), complete cds

gi|215993|gb|M27738|PT6REGA [215993]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 5 nucleotide neighbors)

M38465

Bacteriophage T6 DNA ligase gene, complete cds

gi|215991|gb|M38465|PT6LIG55 [215991]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

V01146

Genome of bacteriophage T7

gi|431187|emb|V01146|T7CG [431187]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,13 MEDLINE links, 60 protein links, 105 nucleotide neighbors, or 1 genome link)

X60322

Bacteriophage alpha3 genes A, B, K, C, D, E, J, F, G, H

gi|14775|emb|X60322|BACALPHA [14775]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 10 protein links, 22 nucleotide neighbors, or 1 genome link)

X13332

Bacteriophage alpha3 DNA for origin of replication

gi|15093|emb|X13332|MLA3ORPL [15093]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

X12611

Bacteriophage alpha3 gene for protein A part., finger domain

gi|15092|emb|X12611|MLA3AFIN [15092]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 6 nucleotide neighbors)

X15721

Bacteriophage alpha3 deletion mutation DNA for the origin region (-ori) of replication

gi|14774|emb|X15721|BA3DMOR9 [14774]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 11 nucleotide neighbors)

X15720

Bacteriophage alpha3 deletion mutant DNA for the origin region (-ori) of replication

gi|14773|emb|X15720|BA3DMOR8 [14773]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

X15719

Bacteriophage alpha3 insertion mutant DNA for the origin region (-ori) of replication

gi|14772|emb|X15719|BA3DMOR7 [14772]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 10 nucleotide neighbors)

X15718

Bacteriophage alpha3 deletion mutation DNA for origin region (-ori) of replication

gi|14771|emb|X15718|BA3DMOR6 [14771]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 11 nucleotide neighbors)

X15717

Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication

gi|14770|emb|X15717|BA3DMOR5 [14770]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 9 nucleotide neighbors)

X15716

Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication

gi|14769|emb|X15716|BA3DMOR4 [14769]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 10 nucleotide neighbors)

X15715

Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication
gi|14768|emb|X15715|BA3DMOR3 [14768]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)

X15714

Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication
gi|14767|emb|X15714|BA3DMOR2 [14767]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)

X15713

Bacteriophage alpha3 deletion mutant DNA for the origin region (-ori) of replication
gi|14766|emb|X15713|BA3DMOR1 [14766]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)

X62059

Bacteriophage alpha3 origin of cDNA synthesis (oriGA)
gi|14763|emb|X62059|AL3ORIGA [14763]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)

X62058

Bacteriophage alpha3 origin of cDNA synthesis (oriAA)
gi|14762|emb|X62058|AL3ORIAA [14762]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)

J02444

Bacteriophage alpha3 origin of DNA replication
gi|166103|gb|J02444|AL3ORI [166103]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors)

M25640

Bacteriophage alpha-3 H protein gene, complete cds
gi|166101|gb|M25640|AL3HP [166101]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 13 nucleotide neighbors)

M10631

Bacteriophage alpha-3 cleavage site for phage phi-X174 gene A protein
gi|166099|gb|M10631|AL3CSA [166099]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

X00774

Bacteriophage alpha-3 gene J sequence
gi|15431|emb|X00774|NCBA3J [15431]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

M25640

Bacteriophage alpha-3 H protein gene, complete cds
gi|166101|gb|M25640|AL3HP [166101]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 13 nucleotide neighbors)

M10631

Bacteriophage alpha-3 cleavage site for phage phi-X174 gene A protein
gi|166099|gb|M10631|AL3CSA [166099]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

227

J02459

Bacteriophage lambda, complete genome

gi|215104|gb|J02459|LAMCG [215104]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,87 MEDLINE links, 67 protein links, 190 nucleotide neighbors, or 1 genome link)

J02482

Bacteriophage phi-X174, complete genome

gi|216019|gb|J02482|PX1CG [216019]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,23 MEDLINE links, 11 protein links, 26 nucleotide neighbors, or 1 genome link)

J02454

Bacteriophage G4, complete genome

gi|215415|gb|J02454|PG4CG [215415]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 11 protein links, 20 nucleotide neighbors, or 1 genome link)

X60323

Bacteriophage phiK complete genome

gi|1478118|emb|X60323|BPHKCG [1478118]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,10 protein links, 18 nucleotide neighbors, or 1 genome link)

L42820

Bacteriophage BF23 tail protein (hrs) gene, complete cds

gi|1048680|gb|L42820|BBFHRS [1048680]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

X54455

Bacteriophage BF23 gene 17 and gene 18

gi|14797|emb|X54455|BF231718G [14797]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 2 nucleotide neighbors)

M37097

Bacteriophage BF23 DNA, right end of terminal repetition

gi|166115|gb|M37097|BBFRIGH [166115]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M37096

Bacteriophage BF23 DNA, left end of terminal repetition

gi|166114|gb|M37096|BBFLEFT [166114]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M37095

Bacteriophage BF23 A2-A3 gene, complete cds, and A1 gene, 5' end

gi|166110|gb|M37095|BBFA2A3 [166110]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 3 protein links, or 1 nucleotide neighbor)

AF056281

Bacteriophage BF23 clone bf23.mac5/6.1, genomic survey sequence

gi|3090930|gb|AF056281|AF056281 [3090930]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056280

Bacteriophage BF23 clone bf23.mac3, genomic survey sequence
gi|3090929|gb|AF056280|AF056280 [3090929]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056279

Bacteriophage BF23 clone bf23.mac18/21.34, genomic survey sequence
gi|3090928|gb|AF056279|AF056279 [3090928]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056278

Bacteriophage BF23 clone bf23.mac16/19.33, genomic survey sequence
gi|3090927|gb|AF056278|AF056278 [3090927]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056277

Bacteriophage BF23 clone bf23.mac16/19-33, genomic survey sequence
gi|3090926|gb|AF056277|AF056277 [3090926]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056276

Bacteriophage BF23 clone bf23.mac12/9-9, genomic survey sequence
gi|3090925|gb|AF056276|AF056276 [3090925]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056275

Bacteriophage BF23 clone bf23.mac11/14-24, genomic survey sequence
gi|3090924|gb|AF056275|AF056275 [3090924]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056274

Bacteriophage BF23 clone bf23.57r64r, genomic survey sequence
gi|3090923|gb|AF056274|AF056274 [3090923]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 3 nucleotide neighbors)

AF056273

Bacteriophage BF23 clone bf23.54fr, genomic survey sequence
gi|3090922|gb|AF056273|AF056273 [3090922]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056272

Bacteriophage BF23 clone bf23.47fr.mac10/7, genomic survey sequence
gi|3090921|gb|AF056272|AF056272 [3090921]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056271

Bacteriophage BF23 clone bf23.23.66r, genomic survey sequence
gi|3090920|gb|AF056271|AF056271 [3090920]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056270

Bacteriophage BF23 clone bf23.23.64f, genomic survey sequence
gi|3090919|gb|AF056270|AF056270 [3090919]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056269

Bacteriophage BF23 clone bf23.23.60r, genomic survey sequence
gi|3090918|gb|AF056269|AF056269 [3090918]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056268

Bacteriophage BF23 clone bf23.23.60f, genomic survey sequence
gi|3090917|gb|AF056268|AF056268 [3090917]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 nucleotide neighbor)

AF056267

Bacteriophage BF23 clone bf23.23.59r, genomic survey sequence
gi|3090916|gb|AF056267|AF056267 [3090916]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056266

Bacteriophage BF23 clone bf23.23.59f, genomic survey sequence
gi|3090915|gb|AF056266|AF056266 [3090915]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056265

Bacteriophage BF23 clone bf23.23.56r, genomic survey sequence
gi|3090914|gb|AF056265|AF056265 [3090914]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056264

Bacteriophage BF23 clone bf23.23.56f, genomic survey sequence
gi|3090913|gb|AF056264|AF056264 [3090913]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056263

Bacteriophage BF23 clone bf23.23.68f55r, genomic survey sequence
gi|3090912|gb|AF056263|AF056263 [3090912]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056262

Bacteriophage BF23 clone bf23.23.43fr.66f, genomic survey sequence
gi|3090911|gb|AF056262|AF056262 [3090911]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056261

Bacteriophage BF23 clone bf23.23.2fr, genomic survey sequence
gi|3090910|gb|AF056261|AF056261 [3090910]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056260

Bacteriophage BF23 clone bf23.23.55.f, genomic survey sequence
gi|3090909|gb|AF056260|AF056260 [3090909]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056259

Bacteriophage BF23 clone bf23.23.53.r, genomic survey sequence
gi|3090908|gb|AF056259|AF056259 [3090908]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056258

Bacteriophage BF23 clone bf23.23.53.f, genomic survey sequence
gi|3090907|gb|AF056258|AF056258 [3090907]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056257

Bacteriophage BF23 clone bf23.23.52.r, genomic survey sequence
gi|3090906|gb|AF056257|AF056257 [3090906]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056256

Bacteriophage BF23 clone bf23.23.52.f, genomic survey sequence
gi|3090905|gb|AF056256|AF056256 [3090905]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056255

Bacteriophage BF23 clone bf23.23.49.r, genomic survey sequence
gi|3090904|gb|AF056255|AF056255 [3090904]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056254

Bacteriophage BF23 clone bf23.23.49.f, genomic survey sequence
gi|3090903|gb|AF056254|AF056254 [3090903]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056253

Bacteriophage BF23 clone bf23.23.48.r, genomic survey sequence
gi|3090902|gb|AF056253|AF056253 [3090902]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056252

Bacteriophage BF23 clone bf23.23.48.f, genomic survey sequence
gi|3090901|gb|AF056252|AF056252 [3090901]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056251

Bacteriophage BF23 clone bf23.23.44.r, genomic survey sequence
gi|3090900|gb|AF056251|AF056251 [3090900]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056250

Bacteriophage BF23 clone bf23.23.41.f, genomic survey sequence
gi|3090899|gb|AF056250|AF056250 [3090899]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056249

Bacteriophage BF23 clone bf23.23.22.a.r, genomic survey sequence
gi|3090898|gb|AF056249|AF056249 [3090898]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056248

Bacteriophage BF23 clone bf23.23.22.a.f, genomic survey sequence
gi|3090897|gb|AF056248|AF056248 [3090897]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056247

Bacteriophage BF23 clone bf23.23.68.r, genomic survey sequence
gi|3090896|gb|AF056247|AF056247 [3090896]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

Z50114

Bacteriophage BF23 DNA for putative tail protein gene
gi|2464952|emb|Z50114|BF23LATE [2464952]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)

D12824

Bacteriophage BF23 genes for minor tail protein gp24 and major tail protein gp25, complete cds
gi|520578|dbj|D12824|BBF2TAIL [520578]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

Z34953

Bacteriophage K3 ip9, ip7 and ip8 genes
gi|535261|emb|Z34953|MYK3IP978 [535261]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

Z35075

Bacteriophage K3 DNA for Ip3 and Ip4
gi|535229|emb|Z35075|MYEORF64K [535229]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

X05560

Bacteriophage K3 gene 38 for receptor recognizing protein
gi|15112|emb|X05560|MYK3G38 [15112]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

X04747

Bacteriophage K3 gene 37 for receptor recognizing protein
gi|15110|emb|X04747|MYK3G37 [15110]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

X01754

Bacteriophage K3 tail fiber gene 36
gi|15108|emb|X01754|MYK3F36 [15108]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

M16812

Bacteriophage K3 'r' lysis gene, complete cds
gi|215503|gb|M16812|PK3LYST [215503]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

L46833

Bacteriophage K3 frd3, frd2 genes, complete cds
gi|951377|gb|L46833|PK3FRD32G [951377]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 2 nucleotide neighbors)

L43613

Bacteriophage K3 fibrin (wac) gene, complete cds
gi|903861|gb|L43613|PK3WAC [903861]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)

X01753

Bacteriophage Ox2 tail fiber gene 36

gi|15122|emb|X01753|MYOX2F36 [15122]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

L43612

Bacteriophage Ox2 fibrin (wac) gene, complete cds

gi|903848|gb|L43612|OX2WAC [903848]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)

Z46880

Bacteriophage OX2 stp gene

gi|599663|emb|Z46880|BPOX2STP [599663]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

X05675

Bacteriophage Ox2 gene 38 for receptor-recognizing protein and flanking regions

gi|15124|emb|X05675|MYOX2G38 [15124]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

M33533

Bacteriophage RB18 translational repressor protein (regA) and Orf43.1, complete cds

gi|216083|gb|M33533|RB18REGA [216083]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

AF033329

Bacteriophage RB18 single-stranded binding protein (gene 32) gene, partial cds, and 5' region

gi|2645788|gb|AF033329|AF033329 [2645788]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 11 nucleotide neighbors)

M86231

Bacteriophage RB69 gene 62, 3'end; RegA (regA) gene, complete cds

gi|215354|gb|M86231|P6962REGA [215354]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

AF033332

Bacteriophage RB69 single-stranded binding protein (gene 32) gene, partial cds, and 5' region

gi|2645794|gb|AF033332|AF033332 [2645794]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 12 nucleotide neighbors)

U34036

Bacteriophage RB69 DNA polymerase (43) gene, complete cds

gi|1237125|gb|U34036|BRU34036 [1237125]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

V01145

Bacteriophage H1 genome fragment Each Thymine given in this sequence represents a HMU-residue (HMU = 5-hydroxymethyluracil)

gi|15557|emb|V01145|PODOH1 [15557]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

X05676

Bacteriophage M1 gene 38 for receptor recognizing protein and flanking regions

gi|15114|emb|X05676|MYM1G38 [15114]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

AF034575

Bacteriophage M1 putative integrase (int) gene, complete cds, and attP region, complete sequence
gi|2662472|gb|AF034575|AF034575 [2662472]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

AF033321

Bacteriophage M1 single-stranded binding protein (gene 32) gene, partial cds, and 5' region
gi|2645772|gb|AF033321|AF033321 [2645772]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 17 nucleotide neighbors)

X55190

Bacteriophage Tu1a 37 and 38 genes for receptor-recognizing proteins 37 and 38 (respectively), partial cds
gi|14860|emb|X55190|BPTUIA [14860]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

AF033334

Bacteriophage Tu1b single-stranded binding protein (gene 32) gene, partial cds, and 5' region
gi|2645798|gb|AF033334|AF033334 [2645798]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 5 nucleotide neighbors)

X55191

Bacteriophage Tu1b 37 gene for receptor-recognizing protein 37 (partial cds), 38 gene for receptor-recognizing protein 38, and t gene (partial cds)
gi|14863|emb|X55191|BPTUIB [14863]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)

X13065

Bacteriophage phi80 early region
gi|14800|emb|X13065|BP80ER [14800]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 8 protein links, or 6 nucleotide neighbors)

D00360

Bacteriophage phi80 cor gene
gi|217782|dbj|D00360|P8080COR [217782]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)

X01639

Bacteriophage phi 80 DNA-fragment with replication origin
gi|15828|emb|X01639|XXPHI80 [15828]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 25 nucleotide neighbors)

X04051

Lambdoid bacteriophage phi 80 int-xis region (integrase-excisionase region)
gi|15770|emb|X04051|STPHI80X [15770]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

X06751

Phage Phi80 DNA for major coat protein
gi|15768|emb|X06751|STPHI80C [15768]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 11 nucleotide neighbors)

X75949

Bacteriophage phi80 DNA for ORF x171.5 and ORF x171.28
gi|458811|emb|X75949|ECORF171B [458811]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 28 nucleotide neighbors)

L40418

Bacteriophage phi-80 gene, complete cds

gi|1019107|gb|L40418|P80A [1019107]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

M24831

Bacteriophage phi-80 Tyr-tRNA gene, 3' end

gi|215363|gb|M24831|P80TGY [215363]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 43 nucleotide neighbors)

M10670

Bacteriophage phi-80 replication origin

gi|215361|gb|M10670|P80ORJ [215361]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

M24825

Bacteriophage phi-80 RNA fragment

gi|215360|gb|M24825|P80M3A [215360]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M11919

Bacteriophage phi-80 cI immunity region encoding the N gene

gi|215358|gb|M11919|P80CI [215358]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

M10891

Bacteriophage phi-80 attP site DNA

gi|215357|gb|M10891|P80ATTIP [215357]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M19473

Bacteriophage 933J (from E.coli) proviral Shiga-like toxin type 1 subunits A and B genes, complete cds

gi|215072|gb|M19473|J93SLTI [215072]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 2 protein links, or 20 nucleotide neighbors)

Y10775

Bacteriophage 933W ileX, stx2A and stx2B genes

gi|1938206|emb|Y10775|BP933ILEX [1938206]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 36 nucleotide neighbors)

X83722

Bacteriophage 933W slt-IIIB gene

gi|1490229|emb|X83722|B933WSLT [1490229]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 20 nucleotide neighbors)

X07865

Bacteriophage 933W slt-II gene for Shiga-like toxin typeII subunit A and B

gi|14892|emb|X07865|BWSLTII [14892]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 29 nucleotide neighbors)

M16625

Bacteriophage H19B (from E.coli) sltIA and sltIB genes encoding Shiga-like toxin I subunits A and B, complete cds

gi|215043|gb|M16625|H19BSLT [215043]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 24 nucleotide neighbors)

235

M17358

Bacteriophage H19B shiga-like toxin-1 (SLT-1) A and B subunit DNA, complete cds

gi|215046|gb|M17358|H19BSLTA [215046]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 20 nucleotide neighbors)

U29728

Bacteriophage N4 single-stranded DNA-binding protein (N4SSB) gene, complete cds

gi|939708|gb|U29728|BNU29728 [939708]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 1 protein link)

J02580

Bacteriophage PA-2 (E.coli porcine strain isolate) Rz gene, 5'end; ORF2, outer membrane porin protein (lc) and ORF1 genes. complete cds

gi|215366|gb|J02580|PA2LC [215366]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 4 nucleotide neighbors)

U32222

Bacteriophage 186, complete sequence

gi|3337249|gb|U32222|B1U32222 [3337249]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 46 protein links, or 5 nucleotide neighbors)

X51522

Bacteriophage P4 complete DNA genome

gi|450916|emb|X51522|MYP4CG [450916]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 13 protein links, 6 nucleotide neighbors, or 1 genome link)

X92588

Bacteriophage 82 orf33, orf151, orf56, orf96, rus, orf45, and Q genes

gi|1051111|emb|X92588|BAC82HOLL [1051111]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,7 protein links, or 1 nucleotide neighbor)

J02803

Bacteriophage 82 antitermination protein (Q) gene, complete cds

gi|215364|gb|J02803|P82Q [215364]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

U02466

Bacteriophage HK022 (cro), (cII) and (O) genes, complete cds, (P) gene, partial cds

gi|407285|gb|U02466|BHU02466 [407285]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 5 protein links, or 1 nucleotide neighbor)

M26291

Bacteriophage D108 regulatory DNA-binding protein (ner) gene, complete cds

gi|166194|gb|M26291|D18NER [166194]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

M11272

Bacteriophage D108 left-end DNA

gi|166193|gb|M11272|D18LEDNA [166193]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M18902

Bacteriophage D108 kil gene encoding a replication protein, 3' end; and containing three ORFs, complete cds

gi|166191|gb|M18902|D18KIL [166191]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

M10191

Bacteriophage D108, left end with Mu A protein binding sites L1 and L2

gi|166190|gb|M10191|D18BSL [166190]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)

J02447

bacteriophage d108 gene a 5' end

gi|166189|gb|J02447|D18AAA [166189]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

V00865

Bacteriophage D108 fragment from genes A and ner (C-terminus of ner and N-terminus of A)

gi|15437|emb|V00865|NCD108 [15437]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

X01914

Bacteriophage IKe gene for DNA binding protein

gi|14957|emb|X01914|INIKEDBP [14957]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

AF064539

Bacteriophage N15, complete genome

gi|3192683|gb|AF064539|AF064539 [3192683]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 60 protein links, 26 nucleotide neighbors, or 1 genome link)

U02303

Bacteriophage If1, complete genome

gi|3676280|gb|U02303|B2U02303 [3676280]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 10 protein links, or 1 genome link)

AF007792

Bacteriophage Mu late morphogenetic region

gi|3551775|gb|AF007792|AF007792 [3551775]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor.)

U24159

Bacteriophage HP1 strain HP1c1, complete genome

gi|1046235|gb|U24159|BHU24159 [1046235]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 6 MEDLINE links, 41 protein links, 8 nucleotide neighbors, or 1 genome link)

Z71579

Bacteriophage S2 type A 5.6 kb DNA fragment

gi|1679806|emb|Z71579|BPHS1ADNA [1679806]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 9 protein links, or 9 nucleotide neighbors)

X53238

Klebsiella sp. bacteriophage K11 gene 1 for RNA polymerase

gi|14984|emb|X53238|KSK11RPO [14984]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

X85010

Bacteriophage A511 ply511 gene

gi|853748|emb|X85010|BPA511PLY [853748]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

U29728

Bacteriophage N4 single-stranded DNA-binding protein (N4SSB) gene, complete cds

gi|939708|gb|U29728|BNU29728 [939708]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 1 protein link)

J02445

bacteriophage bo1 3'-terminal region rna

gi|166152|gb|J02445|BO1TR3 [166152]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 5 nucleotide neighbors)

L06183

Bacteriophage L5 (from *Leuconostoc oenos*) genome

gi|289353|gb|L06183|BL5GENM [289353]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 genome link)

AF074945

Mycoplasma arthritidis bacteriophage MAV1, complete genome

gi|3511243|gb|AF074945|AF074945 [3511243]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,15 protein links, 3 nucleotide neighbors, or 1 genome link)

L13696

Bacteriophage L2 (from *Mycoplasma*), complete genome

gi|289338|gb|L13696|BL2CG [289338]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 14 protein links, or 1 genome link)

X80191

Bacteriophage PP7 mRNA for maturation, coat, lysis and replicase proteins

gi|517237|emb|X80191|BPP7PR [517237]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 1 genome link)

M19377

Bacteriophage Pf3 from *Pseudomonas aeruginosa* (New York strain), complete genome

gi|215380|gb|M19377|PF3COMNY [215380]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 9 protein links, or 5 nucleotide neighbors)

M11912

Bacteriophage Pf3 from *Pseudomonas aeruginosa* (Nijmegen strain), complete genome

gi|215371|gb|M11912|PF3COMN [215371]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 9 protein links, 5 nucleotide neighbors, or 1 genome link)

V00605

Bacteriophage Pf1 gene encoding DNA binding protein

gi|14970|emb|V00605|INOPF1 [14970]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 1 nucleotide neighbor)

L05626

Bacteriophage PR4 capsid protein (P6) gene, complete cds

gi|215735|gb|L05626|PR4P6MAJA [215735]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

D13409

Bacteriophage phiCTX (isolated from *Pseudomonas aeruginosa*) cosR, attP, int genes
gi|217776|dbj|D13409|BPHCOSR [217776]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)

D13408

Bacteriophage phiCTX (isolated from *Pseudomonas aeruginosa*) cosL, ctx genes
gi|217775|dbj|D13408|BPHCOSLCTX [217775]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 3 nucleotide neighbors)

M24832

Bacteriophage f2 coat protein gene, partial cds
gi|166228|gb|M24832|F2CRNACA [166228]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

S72011

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618967|gb|AF017629|AF017629 [2618967]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017628

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618964|gb|AF017628|AF017628 [2618964]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017627

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618961|gb|AF017627|AF017627 [2618961]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017626

Bacteriophage 21 isocitrate dehydrogenase (icd) gene, partial cds; and integrase (int) gene, partial cds
gi|2618958|gb|AF017626|AF017626 [2618958]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 49 nucleotide neighbors)

AF017625

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618955|gb|AF017625|AF017625 [2618955]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017624

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618952|gb|AF017624|AF017624 [2618952]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017623

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618949|gb|AF017623|AF017623 [2618949]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017622

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618946|gb|AF017622|AF017622 [2618946]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017621

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618943|gb|AF017621|AF017621 [2618943]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

D26449

Bacteriophage PS17 FI gene for tail sheath protein (gpFI) and FII gene for tail tube protein (gpFII), complete cds
gi|452162|dbj|D26449|BPSFIFII [452162]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 protein links)

X87627

Bacteriophage D3112 A and B genes
gi|974768|emb|X87627|BPD3112AB [974768]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

U32623

Bacteriophage D3 transcriptional activator CII (cII) gene, complete cds
gi|984852|gb|U32623|BDU32623 [984852]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 1 nucleotide neighbor)

L34781

Bacteriophage phi 11 holin homologue (ORF3) gene, complete cds and peptidoglycan hydrolase (lytA) gene, partial cds
gi|511838|gb|L34781|BPHHOLIN [511838]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 2 nucleotide neighbors)

L14810

Bacteriophage P22 (gp10) gene, complete cds, and (gp26) gene, complete cds
gi|294053|gb|L14810|P22GP1026X [294053]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

X87420

Bacteriophage ES18 genes 24, c2, cro, c1, 18, and oL and oR operators
gi|1143407|emb|X87420|BPES18GEN [1143407]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,5 protein links, or 9 nucleotide neighbors)

L42820

Bacteriophage BF23 tail protein (hrs) gene, complete cds
gi|1048680|gb|L42820|BBFHRS [1048680]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

X14980

Bacteriophage PRD1 XV gene for protein P15 (lytic enzyme)
gi|15802|emb|X14980|TEPRD1XV [15802]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

X06321

Bacteriophage PRD1 gene 8 for DNA terminal protein
gi|15800|emb|X06321|TEPRD18 [15800]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 10 nucleotide neighbors)

X14336

Filamentous Bacteriophage I2-2 genome
gi|14920|emb|X14336|INBI22 [14920]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 9 protein links, 1 nucleotide neighbor, or 1 genome link)

L05001 240

Bacteriophage X glucosyl transferase gene, complete cds
gi|216044|gb|L05001|PXFCLUSYLT [216044]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

M29479

Bacteriophage p4 sid and psu genes partial cds, and delta gene, complete cds gi|215701|
gb|M29479|PP4SDP [215701]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 protein links, or 4 nucleotide neighbors)

SEG_PP4PSUSID

Bacteriophage P4 capsid size determination protein (sid) gene, 5' end
gi|215698|gb|SEG_PP4PSUSID [215698]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

M29650

Bacteriophage P4 polarity suppression protein (psu) gene, complete cds
gi|215697|gb|M29650|PP4PSUSID2 [215697]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

M29651

Bacteriophage P4 capsid size determination protein (sid) gene, 5' end
gi|215696|gb|M29651|PP4PSUSID1 [215696]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

M27748

Bacteriophage P4 gop, beta, and cII genes, complete cds and int gene, 3' end
gi|215691|gb|M27748|PP4GOPBC [215691]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 1 nucleotide neighbor)

K02750

Bacteriophage IKe, complete genome
gi|215061|gb|K02750|IKECG [215061]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, 4 nucleotide neighbors, or 1 genome link)

L40418

Bacteriophage phi-80 gene, complete cds
gi|1019107|gb|L40418|P80A [1019107]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

AF032122

Bacteriophage Sfil integrase (int) gene, partial cds; and bactoprenol glucosyl transferase (bgt), and glucosyl transferase II (grII) genes, complete cds
gi|2465412|gb|AF021347|AF021347 [2465412]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 2 nucleotide neighbors)

M35825

Bacteriophage SF6 fragment D lysozyme gene, complete cds
gi|216105|gb|M35825|SF6LYZ [216105]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)

Z35479

Bacteriophage C16 ip1 gene
gi|534936|emb|Z35479|BC16IP1 [534936]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

X12638

Bacteriophage 21 DNA for gene 2

gi|296141|emb|X12638|B21GENE2 [296141]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

X02501

Bacteriophage 21 DNA for left end sequence with genes 1 and 2

gi|15825|emb|X02501|XXPHA21 [15825]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

M65239

Bacteriophage 21 lysis genes S, R, and Rz, complete cds

gi|215466|gb|M65239|PH2LYSGEN [215466]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

M58702

Bacteriophage 21 late gene regulatory region

gi|215465|gb|M58702|PH2LATEGE [215465]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M81255

Bacteriophage 21 head gene operon

gi|215454|gb|M81255|PH2HEADTL [215454]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 10 protein links, or 4 nucleotide neighbors)

M23775

Bacteriophage 21 glycoprotein 1 gene, complete cds, and glycoprotein gene, 5' end

gi|215451|gb|M23775|PH2GPA [215451]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

M61865

Bacteriophage 21 excisionase (xis), integrase (int) and isocitrate dehydrogenase (icd), complete cds

gi|215448|gb|M61865|PH22XISAA [215448]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 9 nucleotide neighbors)

S72011

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds

gi|2618967|gb|AF017629|AF017629 [2618967]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017628

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds

gi|2618964|gb|AF017628|AF017628 [2618964]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017627

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds

gi|2618961|gb|AF017627|AF017627 [2618961]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017626

Bacteriophage 21 isocitrate dehydrogenase (icd) gene, partial cds; and integrase (int) gene, partial cds

gi|2618958|gb|AF017626|AF017626 [2618958]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 49 nucleotide neighbors)

AF017625

242
Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618955|gb|AF017625|AF017625 [2618955]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017624

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618952|gb|AF017624|AF017624 [2618952]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017623

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618949|gb|AF017623|AF017623 [2618949]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017622

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618946|gb|AF017622|AF017622 [2618946]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017621

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618943|gb|AF017621|AF017621 [2618943]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

M57455

Bacteriophage 42D (clone pDB17) (from *Staphylococcus aureus*) staphylokinase gene, complete cds
gi|215344|gb|M57455|P42STK [215344]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 9 nucleotide neighbors)

Y12633

Bacteriophage 85 DNA, promoter sequence of unknown gene
gi|2058285|emb|Y12633|B85PROM [2058285]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

X98146

Bacteriophage P1 DNA sequence around the Op88 operator
gi|1359513|emb|X98146|BP1OP88OP [1359513]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor)

Y07739

Staphylococcus phage Twort holTW, plyTW genes
gi|2764979|emb|Y07739|BPTWGHOLG [2764979]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2 protein links)

L07580

Bacteriophage phi-11 rinA and rinB genes, required for the activation of *Staphylococcal* phage phi-11 int expression
gi|166160|gb|L07580|BPHRINAB [166160]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

M34832

Bacteriophage phi-11 integrase (int) and excisionase (xis) genes, complete cds
gi|166157|gb|M34832|BPINTXIS [166157]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

- M20394**
Bacteriophage phi-11 S.aureus attachment site (attP)
gi|166156|gb|M20394|BPHATTP [166156]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 nucleotide neighbors)
- X23128**
Bacteriophage phi-13 integrase gene
gi|758228|emb|X82312|PHI13INT [758228]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 3 nucleotide neighbors)
- X61719**
S.aureus phi-13 lysogen right chromosome/bacteriophage DNA junction
gi|46625|emb|X61719|SAP13RJNC [46625]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
- X61718**
S.aureus phi-13 lysogen left chromosomal/bacteriophage DNA junction
gi|46624|emb|X61718|SAP13LJNC [46624]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)
- X61717**
Bacteriophage phi-13 core sequence for attachment
gi|14799|emb|X61717|BP13ATTP [14799]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 3 nucleotide neighbors)
- U01875**
Bacteriophage phi-13 putative regulatory region and integrase (int) gene, partial cds
gi|437118|gb|U01875|U01875 [437118]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, or 4 nucleotide neighbors)
- X67739**
S.aureus Bacteriophage phi-42 attP gene
gi|14809|emb|X67739|BPATTPA [14809]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)
- U01872**
Bacteriophage phi-42 integrase (int) gene, complete cds
gi|437115|gb|U01872|U01872 [437115]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 2 protein links, or 3 nucleotide neighbors)
- X94423**
Staphylococcus aureus bacteriophage phi-42 DNA with ORFs (restriction modification system)
gi|1771597|emb|X94423|SARMS [1771597]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 1 nucleotide neighbor)
- M27965**
Bacteriophage L54a (from S.aureus) int and xis genes, complete cds
gi|215096|gb|M27965|L54INTXIS [215096]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, MEDLINE 1 link, 2 protein links, or 3 nucleotide neighbors)
- U72397**
Bacteriophage 80 alpha holin and amidase genes, complete cds
gi|1763241|gb|U72397|B8U72397 [1763241]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 2 nucleotide neighbors)

AB009866

Bacteriophage phi PVL proviral DNA, complete sequence

gi|3341907|dbj|AB009866|AB009866 [3341907]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,63 protein links, or 1 nucleotide neighbor)

Z47794

Bacteriophage Cp-1 DNA, complete genome

gi|2288892|emb|Z47794|BPCP1XX [2288892]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 28 protein links, 1 nucleotide neighbor, or 1 genome link)

SEG_CP7RSIT

Bacteriophage Cp-7 (S.pneumoniae) 5' inverted terminal repeat

gi|166186|gb|SEG_CP7RSIT [166186]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M11635

Bacteriophage Cp-7 (S.pneumoniae) DNA, 3' inverted terminal repeat

gi|166185|gb|M11635|CP7RSIT2 [166185]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M11636

Bacteriophage Cp-7 (S.pneumoniae) 5' inverted terminal repeat

gi|166184|gb|M11636|CP7RSIT1 [166184]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

SEG_CP5RSIT

Bacteriophage Cp-5 (S.pneumoniae), 5' inverted terminal repeat

gi|166181|gb|SEG_CP5RSIT [166181]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M11633

Bacteriophage Cp-5 (S.pneumoniae) 3' inverted terminal repeat

gi|166180|gb|M11633|CP5RSIT2 [166180]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M11634

Bacteriophage Cp-5 (S.pneumoniae), 5' inverted terminal repeat

gi|166179|gb|M11634|CP5RSIT1 [166179]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M34780

Bacteriophage Cp-9 muramidase (cpl9) gene

gi|166187|gb|M34780|CP9CPL [166187]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

M34652

Bacteriophage HB-3 amidase (hbl) gene, complete cds

gi|215055|gb|M34652|HB3HBLA [215055]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

U64984

Streptococcus pyogenes phage T12 repressor, excisionase (xis), integrase(int) and erythrogenic toxin A precursor (speA) genes, complete cds gi|1877426|gb|U40453|SPU40453 [1877426]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 4 protein links, or 22 nucleotide neighbors)

X12375

Phage CP-T1 (*Vibrio cholerae*) DNA for packaging signal (pac site)

gi|15435|emb|X12375|NCCPPAC [15435]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

AF087814

Vibrio cholerae filamentous bacteriophage fs-2 DNA, complete genome sequence

gi|3702207|dbj|AB002632|AB002632 [3702207]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 9 protein links, or 1 genome link)

D83518

Bacteriophage KVP40 gene for major capsid protein precursor, complete cds

gi|3046858|dbj|D83518|D83518 [3046858]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

AF033322

Bacteriophage PST single-stranded binding protein (gene 32) gene, partial cds, and 5' region

gi|2645774|gb|AF033322|AF033322 [2645774]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 17 nucleotide neighbors)

X94331

Bacteriophage L cro, 24, c2, and c1 genes

gi|1469213|emb|X94331|BLCRO24C [1469213]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 protein links)

U82619

Shigella flexneri bacteriophage V glucosyl transferase (gtr), integrase (int) and excisionase (xis) genes, complete cds

gi|2465470|gb|U82619|SFU82619 [2465470]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 8 protein links, or 1 nucleotide neighbor)

246
Table 12NCBI *Entrez* Nucleotide QUERY

Key words: bacteriophage and lysis

56 citations found (all selected)

AJ011581

Bacteriophage PS119 lysis genes 13, 19, 15, and packaging gene 3, complete cds
gil3676084|embl|AJ011581|BPS011581 [3676084]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 protein links, or 1 nucleotide neighbor)

AJ011580

Bacteriophage PS34 lysis genes 13, 19, 15, antiterminator gene 23, and packaging gene 3, complete cds
gil3676078|embl|AJ011580|BPS011580 [3676078]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 protein links, or 2 nucleotide neighbors)

AJ011579

Bacteriophage PS3 lysis genes 13, 19, 15, and packaging gene 3
gil3676073|embl|AJ011579|BPS011579 [3676073]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 protein links, or 1 nucleotide neighbor)

AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cl protein (cl), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds
gil2668751|gb|AF034975 [2668751]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

U37314

Bacteriophage lambda Rz1 protein precursor (Rz1) gene, complete cds
gil1017780|gb|U37314|BLU37314 [1017780]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 1 protein link, or 9 nucleotide neighbors)

U00005

E. coli hflA locus encoding the hflX, hflK and hflC genes, hsq gene, complete cds; miaA gene, partial cds
gil436153|gb|U00005|ECOHLA [436153]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 MEDLINE links, 1 protein link, or 9 nucleotide neighbors)

links, 5 protein links, or 8 nucleotide neighbors)

U32222

Bacteriophage 186, complete sequence
gil3337249|gb|U32222|B1U32222 [3337249]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE
links, 46 protein links, or 5 nucleotide neighbors)

AF064539

Bacteriophage N15, complete genome
gil3192683|gb|AF064539|AF064539 [3192683]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE
links, 60 protein links, 26 nucleotide neighbors, or 1 genome link)

AF063097

Bacteriophage P2, complete genome
gil3139086|gb|AF063097|AF063097 [3139086]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,21 MEDLINE
links, 42 protein links, 3 nucleotide neighbors, or 1 genome link)

Z97974

Bacteriophage phiadh lys, hol, intG, rad,and tec genes
gil2707950|emb|Z97974|BPHIADH [2707950]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE
links, 9 protein links, or 1 nucleotide neighbor)

AF059243

Bacteriophage NL95, complete genome
gil3088545|gb|AF059243|AF059243 [3088545]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE
links, 4 protein links, 3 nucleotide neighbors, or 1 genome link)

AF052431

Bacteriophage M11 A-protein, coat protein, A1-protein, and replicase
genes, complete cds
gil2981208|gb|AF052431| [2981208]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE
links, 4 protein links, or 8 nucleotide neighbors)

Y07739

Staphylococcus phage Twort holTW, plyTW genes
gil2764979|emb|Y07739|BPTWGHOLG [2764979]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2
protein links)

X94331

Bacteriophage L cro, 24, c2, and c1 genes
gil1469213|emblX94331|BLCRO24C [1469213]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 protein links)

X78410

Bacteriophage phiadh holin and lysin genes
gil793848|emblX78410|LGHOLLYS [793848]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

X99260

Bacteriophage B103 genomic sequence
gil1429229|emblX99260|BB103G [1429229]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 17 protein links, or 12 nucleotide neighbors)

AJ000741

Bacteriophage P1 darA operon
gil2462938|emblAJ000741|BPAJ7641 [2462938]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, or 31 nucleotide neighbors)

X87420

Bacteriophage ES18 genes 24, c2, cro, c1, 18, and oL and oR operators
gil1143407|emblX87420|BPES18GEN [1143407]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 protein links, or 9 nucleotide neighbors)

L35561

Bacteriophage phi-105 ORFs 1-3
gil532218|gbL35561|PH5ORFHTR [532218]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 protein links)

D10027

Group II RNA coliphage GA genome
gil217784|dbjD10027|PGAXX [217784]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, 5 nucleotide neighbors, or 1 genome link)

V01128

Bacteriophage phi-X174 (cs70 mutation) complete genome
gil15535|emblV01128|PHIX174 [15535]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 MEDLINE links, 11 protein links, or 26 nucleotide neighbors)

S81763

coat gene...replicase gene [bacteriophage KU1, host=Escherichia coli, group II RNA phage, Genomic RNA, 3 genes, 120 nt]
gil1438766|gb|S81763|S81763 [1438766]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

U38906

Bacteriophage r1t integrase, repressor protein (rro), dUTPase, holin and lysin genes, complete cds
gil1353517|gb|U38906|BRU38906 [1353517]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 50 protein links, or 3 nucleotide neighbors)

X91149

Bacteriophage phi-C31 DNA cos region
gil1107473|emb|X91149|APHIC31C [1107473]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 6 protein links, or 1 nucleotide neighbor)

V00642

phage MS2 genome
gil15081|emb|V00642|LEMS2X [15081]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 8 MEDLINE links, 4 protein links, or 20 nucleotide neighbors)

V01146

Genome of bacteriophage T7
gil431187|emb|V01146|T7CG [431187]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 13 MEDLINE links, 60 protein links, 105 nucleotide neighbors, or 1 genome link)

X78401

Bacteriophage P22 right operon, orf 48, replication genes 18 and 12, nin region genes, ninG phosphatase, late control gene 23, orf 60, complete cds, late control region, start of lysis gene 13
gil512343|emb|X78401|POP22NIN [512343]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 13 protein links, or 4 nucleotide neighbors)

Y00408

Bacteriophage T4 gene t for lysis protein
gil15368|emb|Y00408|MYT4T [15368]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

Z26590

Bacteriophage mv4 lysA and lysB genes
gi1410500|embl|Z26590|MV4LYSAB [410500]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 4 protein links)

X07809

Phage phiX174 lysis (E) gene upstream region
gi15094|embl|X07809|MIPHIXE [15094]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 4 nucleotide neighbors)

Z34528

Lactococcal bacteriophage c2 lysin gene
gi1506455|embl|Z34528|LBC2LYSIN [506455]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

X15031

Bacteriophage fr RNA genome
gi15071|embl|X15031|LEBFRX [15071]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, 9 nucleotide neighbors, or 1 genome link)

X80191

Bacteriophage PP7 mRNA for maturation, coat, lysis and replicase proteins
gi1517237|embl|X80191|BPP7PR [517237]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 1 genome link)

X85010

Bacteriophage A511 ply511 gene
gi1853748|embl|X85010|BPA511PLY [853748]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

X85009

Bacteriophage A500 hol500 and ply500 genes
gi1853744|embl|X85009|BPA500PLY [853744]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 4 nucleotide neighbors)

X85008

Bacteriophage A118 hol118 and ply118 genes
gi1853740|embl|X85008|BPA118PLY [853740]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

251

Z35638

Bacteriophage phi-X174 genes for lysis protein and beta-lactamase
gil520996|embl|Z35638|BPLYSPR [520996]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 2 protein links, or 516 nucleotide neighbors)

J02459

Bacteriophage lambda, complete genome
gil215104|gb|J02459|LAMCG [215104]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,87 MEDLINE
links, 67 protein links, 190 nucleotide neighbors, or 1 genome link)

X87674

Bacteriophage P1 lydA & lydB genes
gil974763|embl|X87674|BACP1LYD [974763]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 2 protein links, or 2 nucleotide neighbors)

X87673

Bacteriophage P1 gene 17
gil974761|embl|X87673|BACP117 [974761]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 1 protein link, or 1 nucleotide neighbor)

M14784

Bacteriophage T3 strain amNG220B right end, tail fiber protein, lysis
protein and DNA packaging proteins, complete cds
gil215810|gb|M14784|PT3RE [215810]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 9 protein links, or 10 nucleotide neighbors)

M11813

Bacteriophage PZA (from B.subtilis), complete genome
gil216046|gb|M11813|PZACG [216046]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE
links, 27 protein links, 17 nucleotide neighbors, or 1 genome link)

M16812

Bacteriophage K3 't' lysis gene, complete cds
gil215503|gb|M16812|PK3LYST [215503]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 1 protein link, or 4 nucleotide neighbors)

J04356

Bacteriophage P22 proteins 15 (complete cds), and 19 (3' end) genes
gil215265|gb|J04356|P2215P [215265]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

J04343

Bacteriophage JP34 coat and lysis protein genes, complete cds, and replicase protein gene, 5' end
gil215076|gb|J04343|JP3COLY [215076]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

J02482

Bacteriophage phi-X174, complete genome
gil216019|gb|J02482|PX1CG [216019]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,23 MEDLINE links, 11 protein links, 26 nucleotide neighbors, or 1 genome link)

M99441

Bacteriophage T4 anti-sigma 70 protein (asiA) gene, complete cds and lysis protein, 3' end
gil215820|gb|M99441|PT4ASIA [215820]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 2 protein links, or 2 nucleotide neighbors)

M65239

Bacteriophage 21 lysis genes S, R, and Rz, complete cds
gil215466|gb|M65239|PH2LYSGEN [215466]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

M10637

Phage G4 D/E overlapping gene system, encoding D (morphogenetic) and E (lysis) proteins
gil215427|gb|M10637|PG4DE [215427]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors)

J02454

Bacteriophage G4, complete genome
gil215415|gb|J02454|PG4CG [215415]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 11 protein links, 20 nucleotide neighbors, or 1 genome link)

J02580

Bacteriophage PA-2 (E.coli porcine strain isolate) Rz gene, 5'end; ORF2, outer membrane porin protein (Ic) and ORF1 genes, complete cds
gil215366|gb|J02580|PA2LC [215366]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 4 nucleotide neighbors)

M14782

Bacillus phage phi-29 head morphogenesis, major head protein, head fiber protein, tail protein, upper collar protein, lower collar protein, pre-neck appendage protein, morphogenesis(13), lysis, morphogenesis(15), encapsidation genes, complete cds
gil215323|gb|M14782|P29LATE2 [215323]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 11 protein links, or 11 nucleotide neighbors)

M10997

Bacteriophage P22 lysis genes 13 and 19, complete cds
gil215262|gb|M10997|P221319 [215262]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

J02467

Bacteriophage MS2, complete genome
gil215232|gb|J02467|MS2CG [215232]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,8 MEDLINE links, 4 protein links, 20 nucleotide neighbors, or 1 genome link)

M14035

Bacteriophage lambda lysis S gene with mutations leading to nonlethality of S in the plasmid pRG1
gil215180|gb|M14035|LAMLYS [215180]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 14 nucleotide neighbors)

U04309

Bacteriophage phi-LC3 putative holin (lysA) gene and putative murein hydrolase (lysB) gene, complete cds
gil530796|gb|U04309|BPU04309 [530796]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

Table 13

NCBI *Entrez* Nucleotide QUERY**Key word: holin****51 citations found (all selected)**

AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cI protein (cI), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds
gil2668751|gb|AF034975| [2668751]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

U52961

Staphylococcus aureus holin-like protein LrgA (lrgA) and LrgB (lrgB) genes, complete cds
gil1841516|gb|U52961|SAU52961 [1841516]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

U28154

Haemophilus somnus cryptic prophage genes, capsid scaffolding protein gene, partial cds, major capsid protein precursor, endonuclease, capsid completion protein, tail synthesis proteins, holin, and lysozyme genes, complete cds
gil1765928|gb|U28154|HSU28154 [1765928]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 13 protein links)

AF032122

Streptococcus thermophilus bacteriophage Sfi19 central region of genome
gil2935682|gb|AF032122| [2935682]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 14 protein links, or 2 nucleotide neighbors)

AF032121

Streptococcus thermophilus bacteriophage Sfi21 central region of genome
gil2935667|gb|AF032121|AF032121 [2935667]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 14 protein links, or 2 nucleotide neighbors)

AF021803

Bacillus subtilis 168 prophage SPbeta N-acetylmuramoyl-L-alanine amidase (blyA), holin-like protein (bhlA), holin-like protein (bhlB), and yolK genes, complete cds; and yolJ gene, partial cds
gi12997594|gb|AF021803|AF021803 [2997594]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 5 protein links, or 1 nucleotide neighbor)

AF057033

Streptococcus thermophilus bacteriophage sf11 gp502 (orf502), gp284 (orf284), gp129 (orf129), gp193 (orf193), gp119 (orf119), gp348 (orf348), gp53 (orf53), gp113 (orf113), gp104 (orf104), gp114 (orf114), gp128 (orf128), gp168 (orf168), gp117 (orf117), gp105 (orf105), putative minor tail protein (orf1510), putative minor structural protein (orf512), putative minor structural protein (orf1000), gp373 (orf373), gp57 (orf57), putative anti-receptor (orf695), putative minor structural protein (orf669), gp149 (orf149), putative holin (orf141), putative holin (orf87), and lysin (orf288) genes, complete cds
gi13320432|gb|AF057033|AF057033 [3320432]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,25 protein links, or 1 nucleotide neighbor)

U32222

Bacteriophage 186, complete sequence
gi13337249|gb|U32222|B1U32222 [3337249]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 46 protein links, or 5 nucleotide neighbors)

AB009866

Bacteriophage phi PVL proviral DNA, complete sequence
gi13341907|dbj|AB009866|AB009866 [3341907]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,63 protein links, or 1 nucleotide neighbor)

AF009630

Bacteriophage bIL170, complete genome
gi13282260|gb|AF009630|AF009630 [3282260]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,63 protein links, 3 nucleotide neighbors, or 1 genome link)

AF064539

Bacteriophage N15, complete genome

gil3192683|gb|AF064539|AF064539 [3192683]
 (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE
 links, 60 protein links, 26 nucleotide neighbors, or 1 genome link)

AF063097

Bacteriophage P2, complete genome
 gil3139086|gb|AF063097|AF063097 [3139086]
 (View GenBank report,FASTA report,ASN.1 report,Graphical view,21 MEDLINE
 links, 42 protein links, 3 nucleotide neighbors, or 1 genome link)

Z97974

Bacteriophage phiadh lys, hol, intG, rad, and tec genes
 gil2707950|embl|Z97974|BPHIADH [2707950]
 (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE
 links, 9 protein links, or 1 nucleotide neighbor)

X95646

Streptococcus thermophilus bacteriophage Sfi21 DNA; lysogeny module,
 8141 bp
 gil2292747|embl|X95646|BSFI21LYS [2292747]
 (View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE
 links, 19 protein links, or 3 nucleotide neighbors)

SEG_LLHLYSINO

Bacteriophage LL-H structural protein gene, partial cds; minor
 structural protein gp61 (g57), unknown protein, unknown protein,
 structural protein (g20), unknown protein, unknown protein, major capsid
 protein (g34), main tail protein gp19 (g17), holin (hol), muramidase
 (mur), unknown protein, unknown protein, unknown protein, unknown
 protein, unknown protein, and unknown protein genes, complete cds;
 unknown protein gene, partial cds; and unknown protein, unknown protein,
 unknown protein, unknown protein, unknown protein, minor structural
 protein gp75 (g70), minor structural protein gp89 (g88), minor
 structural protein gp58 (g71), unknown protein, unknown protein, unknown
 protein, and unknown protein genes, complete cds
 gil1004337|gb|SEG_LLHLYSINO [1004337]
 (View GenBank report,FASTA report,ASN.1 report,Graphical view,4 MEDLINE
 links, 31 protein links, or 1 nucleotide neighbor)

M96254

Bacteriophage LL-H holin (hol), muramidase (mur), and unknown protein
 genes, complete cds
 gil1004336|gb|M96254|LLHLYSINO3 [1004336]
 (View GenBank report,FASTA report,ASN.1 report, or Graphical view)

Y07740

Staphylococcus phage 187 ply187 and hol187 genes
gil2764982|embl|Y07740|BP187PLYH [2764982]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2 protein links)

U88974

Streptococcus thermophilus bacteriophage 01205 DNA sequence
gil2444080|gb|U88974| [2444080]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 57 protein links, or 6 nucleotide neighbors)

Z99117

Bacillus subtilis complete genome (section 14 of 21): from 2599451 to 2812870
gil2634966|embl|Z99117|BSUB0014 [2634966]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 233 protein links, 51 nucleotide neighbors, or 1 genome link)

Z99115

Bacillus subtilis complete genome (section 12 of 21): from 2195541 to 2409220
gil2634478|embl|Z99115|BSUB0012 [2634478]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 244 protein links, 64 nucleotide neighbors, or 1 genome link)

Z99110

Bacillus subtilis complete genome (section 7 of 21): from 1194391 to 1411140
gil2633472|embl|Z99110|BSUB0007 [2633472]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 226 protein links, 31 nucleotide neighbors, or 1 genome link)

X78410

Bacteriophage phiadh holin and lysin genes
gil793848|embl|X78410|LGHOLLYS [793848]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

Z93946

258

Bacteriophage Dp-1 dph and pal genes and 5 open reading frames
gil1934760|embl|Z93946|BPDP1ORFS [1934760]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 6
protein links)

AF011378

Bacteriophage sk1 complete genome
gil2392824|gb|AF011378|AF011378 [2392824]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,54 protein
links, 2 nucleotide neighbors, or 1 genome link)

Z47794

Bacteriophage Cp-1 DNA, complete genome
gil2288892|embl|Z47794|BPCP1XX [2288892]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE
links, 28 protein links, 1 nucleotide neighbor, or 1 genome link)

L35561

Bacteriophage phi-105 ORFs 1-3
gil532218|gb|L35561|PH5ORFHTR [532218]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, or 3 protein links)

D49712

Bacillus licheniformis DNA for ORFs, xpaL2 homologous protein and xpaL1
homologous protein, complete and partial cds
gil1514423|dbj|D49712|D49712 [1514423]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE
links, or 4 protein links)

X90511

Lactobacillus bacteriophage phig1e DNA for Rorf162, Holin, Lysin, and
Rorf175 genes
gil1926386|embl|X90511|LBPHIHOL [1926386]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,4 protein
links, or 1 nucleotide neighbor)

X98106

Lactobacillus bacteriophage phig1e complete genomic DNA
gil1926320|embl|X98106|LBPHIG1E [1926320]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE

link, 50 protein links, or 4 nucleotide neighbors)²⁵⁹

U72397

Bacteriophage 80 alpha holin and amidase genes, complete cds
gil1763241|gb|U72397|B8U72397 [1763241]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 2 nucleotide neighbors)

U38906

Bacteriophage rlt integrase, repressor protein (rro), dUTPase, holin and lysin genes, complete cds
gil1353517|gb|U38906|BRU38906 [1353517]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 50 protein links, or 3 nucleotide neighbors)

X91149

Bacteriophage phi-C31 DNA cos region
gil1107473|embl|X91149|APHIC31C [1107473]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 6 protein links, or 1 nucleotide neighbor)

U24159

Bacteriophage HP1 strain HP1c1, complete genome
gil1046235|gb|U24159|BHU24159 [1046235]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 41 protein links, 8 nucleotide neighbors, or 1 genome link)

Z26590

Bacteriophage mv4 lysA and lysB genes
gil410500|embl|Z26590|MV4LYSAB [410500]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 4 protein links)

Z70177

B.subtilis DNA (28 kb PBSX/skin element region)
gil1225934|embl|Z70177|BSPBSXSE [1225934]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,32 protein links, or 4 nucleotide neighbors)

Z36941

²⁶⁰
B.subtilis defective prophage PBSX xhlA, xhlB, and xylA genes
gil535793|embl|Z36941|BSPBSXXHL [535793]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,4 protein
links, or 5 nucleotide neighbors)

X89234

L.innocua DNA for phagelysin and holin gene
gil1134844|embl|X89234|LICPLYHOL [1134844]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 2 protein links, or 4 nucleotide neighbors)

X85010

Bacteriophage A511 ply511 gene
gil853748|embl|X85010|BPA511PLY [853748]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 3 protein links, or 1 nucleotide neighbor)

X85009

Bacteriophage A500 hol500 and ply500 genes
gil853744|embl|X85009|BPA500PLY [853744]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 3 protein links, or 4 nucleotide neighbors)

X85008

Bacteriophage A118 hol118 and ply118 genes
gil853740|embl|X85008|BPA118PLY [853740]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 3 protein links, or 1 nucleotide neighbor)

L34781

Bacteriophage phi 11 holin homologue (ORF3) gene, complete cds and
peptidoglycan hydrolase (lytA) gene, partial cds
gil511838|gb|L34781|BPHHOLIN [511838]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 4 protein links, or 2 nucleotide neighbors)

U11698

Serratia marcescens SM6 extracellular secretory protein (nucE), putative
phage lysozyme (nucD), and transcriptional activator (nucC) genes,
complete cds
gil509550|gb|U11698|SMU11698 [509550]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE

link, 3 protein links, or 1 nucleotide neighbor)

U31763

Serratia marcescens phage-holin analog protein (regA), putative phage lysozyme (regB), and transcriptional activator (regC) genes, complete cds

gi|965068|gb|U31763|SMU31763 [965068]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

X87674

Bacteriophage P1 lydA & lydB genes

gi|974763|embl|X87674|BACP1LYD [974763]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

L48605

Bacteriophage c2 complete genome

gi|1146276|gb|L48605|C2PVCG [1146276]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 39 protein links, 3 nucleotide neighbors, or 1 genome link)

L33769

Bacteriophage bIL67 DNA polymerase subunit (ORF3-5), essential recombination protein (ORF13), lysin (ORF24), minor tail protein (ORF31), terminase subunit (ORF32), holin (ORF37), unknown protein (ORF 1-2,6-12,14-23,25-30,33-36), complete genome

gi|522252|gb|L33769|L67CG [522252]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 37 protein links, 2 nucleotide neighbors, or 1 genome link)

L31348

Bacteriophage Tuc2009 integrase (int) gene, complete cds; lysin (lys) gene, 3' end

gi|508612|gb|L31348|TU2INT [508612]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 3 protein links, or 3 nucleotide neighbors)

L31364

Bacteriophage Tuc2009 holin (S) gene, complete cds; lysin (lys) gene, complete cds

gi|496281|gb|L31364|TU2SLYS [496281]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

L31366

Bacteriophage Tuc2009 structural protein (mp2) gene, complete cds
gil496278|gb|L31366|TU2MP2A [496278]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

L31365

Bacteriophage Tuc2009 structural protein (mp1) gene, complete cds
gil496276|gb|L31365|TU2MP1A [496276]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

U04309

Bacteriophage phi-LC3 putative holin (lysA) gene and putative murein
hydrolase (lysB) gene, complete cds
gil530796|gb|U04309|BPU04309 [530796]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

Table 14

NCBI *Entrez* Nucleotide QUERY**Key word: bacteriophage and kil****5 citations found (all selected)****AF034975**

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cI protein (cI), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds
gil2668751|gb|AF034975| [2668751]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

X15637

Bacteriophage P22 P(L) operon encompassing ral, 17, kil and arf genes
gil15646|emb|X15637|POP22PL [15646]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 7 protein links, or 2 nucleotide neighbors)

J02459

Bacteriophage lambda, complete genome
gil215104|gb|J02459|LAMCG [215104]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,87 MEDLINE links, 67 protein links, 190 nucleotide neighbors, or 1 genome link)

M64097

Bacteriophage Mu left end
gil215543|gb|M64097|PMULEFTEN [215543]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 39 protein links, or 15 nucleotide neighbors)

M18902

Bacteriophage D108 kil gene encoding a replication protein, 3' end; and containing three ORFs, complete cds
gil166191|gb|M18902|D18KIL [166191]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

Table 15

U77328	V01282	U11787	U93688	A47599	D21131	U76864	U38428
AF151117	AF121672	U11786	U93687	A47598	D30690	U76863	U66665
AF151218	AF072726	U11785	AJ224764	A47597	D14711	U76862	U66664
AF146368	AF115379	U11784	AF064774	A47596	D90119	U76861	U66663
AF144661	AF034153	U11783	AF064773	A47595	D00730	U76860	X87104
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AF060189	Y16431	AF027155	Y14051	AF003592	Z84573	Y09570	X62282
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X81586	X02166	A12903	X61718	U10927	M18264	M14372	M62650
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X62992	X68417	A12899	X67740	U20782	M20270	M36694	M21854
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X14827	X17679	A12897	U02910	L37597	M33479	M12715	L14020
X13404	X63072	A12896	AH003349	L36472	M94061	J04151	M81736
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Z16422	A19943	A04512	U35773	L23109	M97169	L13377	L22565
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Z33407	A19940	X53818	U36379	J02615	M25257	L13374	L19298
Z33406	A19939	M20129	U06451	M18970	M25256	M17348	M80252
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Z33404	A19937	L43082	U20794	M21136	M25254	M17347	
X75439	A19936	X03216	L25426	M10501	M25253	M28364	
X62587	A17958	X70648	M86227	AH000935	M25252	M21319	

Table 16

Phage 44AHJD complete genome sequence. 16668 nucleotides.

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351    agaacaatc gaagaagaaa atgaaaacaa attagaacct actacaacag atgaagatag ttcgaaattt
421    gaccctgttg tattagaaca acgtattgct tcattagaac aacaagtgc tactttttta tcttcacaaa
491    tgcaacaacc acaacaagta caacaacac aatcagatgt aacagaatca aacaagaag ataacgacta
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631    catgctgtct atgatgggtt catcatatga agattcaaga ttaaataaac gaacagaatt aaatgaaac
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1051   acaatatagt gaagaatcag tgattatgga cacagtacca attaacatgg acttatctaa aatcgaggaa
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1891   attccataac cctgaatttg atgaagttac acactggatt cattactatt catttaaaag cattagtcca
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9871 gaattacat gggcgtatat gtcacaacta catggtaaaa gacaacctac cgacgacggt caaatcaaca
9941 acggtcagc tgatgtgtac gtctataaaa agttaggtgc aaaaacaaca cataatccaa cagtagggtt
10011 tgggttctct agtaaacac catacttaca agcaactgca tatggtattg gtcacacagg tgtgtgtga

10081 gcagtttttg aagatgggtc gtttttagtt gcaaaactata atgtaccacc atatgttgca ccatcacgtg
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 10221 tgctttaatta actatgctat aatgaacaca tgctagtaat gctagtaaat aaaatacaaa acataaatcaa
 10291 ttttcgtaca catttttcat gttatctcaa aaagaaaagg agactgttat ttaacagtt gcttttttt
 10361 atttcacatcat gttcacgttt taatatatgc aaatcagatt tgttatgtac tgaacgttca acgtgaaata
 10431 agtcgtttaag tgaaaatgaa ccgatgtcac tttcaatata aagaatatca tcaaatggac tatggctgaa
 10501 attttctcta gcgtctttta atataaatc acgtttcata ttaagttcat cagtaaaata ttcacatcat
 10571 acattaccac atacaatttc agtttttagac ggatatatcg atattgtacc ttgtctatta tagatacttt
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 11691 agaattgattt aatatagtc taaaaattca tatcatggaa atgataatgt gtataagata ttttaatatc
 11761 ttgatatttg ttgagtaact gaaaacgtgt catttcatta ttcaagtaag attccataat attcaatgaa
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 12111 tatcaataat attaaattta aaaccattta aaacattgt taaatctaaa ttgattgaag atttaacacg
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 12321 aaaaatgatt atcgtattta ttacagttat gtgcaatcat gataatatct gtttttgatt ttgtgtattg
 12391 atcacgtctt ttcatatcag tataaaatgc gtcataaaaa gattcgaaac tcggaaatc ttcaacatca
 12461 tcttctcgag atttttaata tatattttcg cgtgtaatat tatcaaaata acgcatgggt tctttaagta
 12531 gtcgcgttaac tttattgtac gctaatgttt ctatatccca gtataaaatc attcgcagtt catgtttatg
 12601 atattgcatg cattctagta atcccataat cttacacacc ttttataagc catattgttt cattagatac
 12671 tttttctgat tctctatata gttatcttgc tatatttttt cttttcttca aaactcactc atctttttct
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 12811 cagctaatatt tttagtagaag cattgtcaaa atgtaaattg cttggattgt agtaataacg ttccatgttt
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 13161 tggtaacgtt ttcatcagc taatcgtttc gtcgcatttc taaaaaaatg tttttgtaaa gtcttgatgt
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 14211 agcattttta tatgatacgt tttcttcttt aggaaaaatg ggcagatgtg caaaatgttt ccatgtgtca
 14281 atgtacgcct cttgtaaaac tttatcatca aatttataat taacattact aaaaatcattt aaaaaataat
 14351 ctttttcttg ctcttttcta gcttctcttt ctttttccca tctatccatt tcagacgtat gtctaaccac
 14421 tgttatcaac ctccatataa agcataaata accattaaaa agataatata gaatataatc aatgtagtga
 14491 ataaaacacc aaatgacacg cgtatatgca gtgtcataag tatgataagt gtaattaaaa atgctaaaaa
 14561 gaaaacaatg gctatgttta ataggttatt catggttcaat cactttccca ttatcgtata tgactttgtt
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 14701 tttataaaaa tttctcttat taattcatta cttaataaat tcttataata aaatacaagt atattaaaaa
 14771 catgtttttt aatatcaatg tcgatataca acgtataata ctcttttcca atttcaaaat catcatattg
 14841 tttgtcaaac tcaatatata catcacccat atttattttt actatacat ttttattaga ttgagtaaat
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 15051 ttatattcat aatatgaata tacaacttta ccgtcatata aatcttcaaa cattgagatt tgatgtggaa
 15121 aatgtctctt aatctcatcg caataataa ataccgtttt gtatttacgt tccattttaa cactcataa
 15191 aaaaatgggg ataagtatcc cctatgaaat tgtattaaaa tgatacttga ccaaaattga ttgagtaacc
 15261 tttttgacct tttttgtttt catattcata aattgtgaat tgaacttctc cagcattgat aatgtcaaca
 15331 acgtctctcat ctgctctcat tttcttaatt aattctgtta agtgggtcgg taagtttacg ttatagtcac

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15401 cagtgacgat aacaccttgt tcaccgaatt ttgattcttt gtttgtgaat aatgctctaa cgatatactc
15471 ttttttcata ccgtattttt ctactaatcc tgatagtttg ataaattctc tttctttttc ctcaaattca
15541 aatctcgccta atgtgttttg gtgtcttgat aaaatatctt ttacgtttgt cattttattt ctctcttat
15611 ttaaattatt tgctttctgc aattgcgatt tgtagtaaat cattgtaata aacttgaatt gttttcgttg
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15751 ctcgtgaagt ggtaaaaatt cctcaatgta ttcattatca tcatctaagt aatgaagtat ataacccttg
15821 acacgtaagg taacaatgtc gtcaactttc attattatat cactcctttc taaaaaacgt aaacgttata
15891 cgtttcataa aatcctttat gcataattcca ttgttctatt gggtcacac cagcaatata agacaatatt
15961 gattctggtt tagtttcgtt gtttagttca tcatttaaga attgaacaac agaactatta tagtttaata
16031 atagtgttg gcaagccgat aataagttaa ttgcattgtc aaatgtataa gctggattcc attgaatcag
16101 tttattgaat agttgcaaca tttcagtata ggcttgcct tttcttctg gtgcattatc aacattaacc
16171 attattatca cttcctaata aagttgaaat tacgcgtaaa acagaattat gatttaaatc ttcaatttca
16241 tcaatgtcaa catcataaaa tgaaatttca ttttctgttc tatcaataa cyctatacat aaacttccat
16311 tcttaaaacg aaaaacatgc ttcaactcaa tgttttttgt ttcatcttcc atttttgta ctcttctgtt
16381 tgattacata cttagtatag caaacgttta aaagttttgt caatagtttt tcttaaaaaa gtttaataa
16451 ttttaaaact actatttaat agaagaaata agattttaag ttcaaatcat aattttgaat aaaagtcaat
16521 agatacataa attttgtatt tgatgaatat gtaatagggt agataagttg gtttaagtgt tgcacagtat
16591 ttttaagttt agtaaagaaa tgataagtaa atttataagt tttgatttgt ataatcgttt attttaaacc
16661 ggtgggggt
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Table 17

Phage 44AHJD ORFs list

nb	Name	Frame	Position	Size (a.a.)	Key words
1	44AHJDORF001	-1	10342..12627	761	DNA polymerase;
2	44AHJDORF002	3	3789..5732	647	Techoic acid; Staph;
3	44AHJDORF003	2	6626..8389	587	Tall;
4	44AHJDORF004	1	8764..10227	487	Serine protease motif;
5	44AHJDORF005	-1	12643..13890	415	
6	44AHJDORF006	2	803..2029	408	
7	44AHJDORF007	1	2044..3027	327	Upper collar;
8	44AHJDORF008	2	3020..3775	251	Lower collar;
9	44AHJDORF009	2	5744..6496	250	Amidase; Staph;
10	44AHJDORF010	-2	13938..14420	160	
11	44AHJDORF012	3	8391..8813	140	Holin;
12	44AHJDORF013	-2	14586..14996	136	
13	44AHJDORF113	1	199..600	133	
14	44AHJDORF011	-2	15225..15593	122	
15	44AHJDORF114	-2	15870..16172	100	
16	44AHJDORF014	3	6243..6521	92	
17	44AHJDORF015	1	15403..15645	80	
18	44AHJDORF016	-1	15616..15852	78	
19	44AHJDORF017	-2	10536..10757	73	
20	44AHJDORF018	-1	886..1098	70	
21	44AHJDORF019	-2	9630..9836	68	
22	44AHJDORF121	-1	16165..16362	65	
23	44AHJDORF020	2	13865..14053	62	
24	44AHJDORF123	2	614..796	60	
25	44AHJDORF021	-2	5634..5816	60	
26	44AHJDORF023	-2	6315..6494	59	
27	44AHJDORF024	1	14275..14451	58	
28	44AHJDORF025	-3	14999..15175	58	
29	44AHJDORF026	-3	14426..14593	55	
30	44AHJDORF027	1	12916..13080	54	
31	44AHJDORF029	-1	15019..15183	54	
32	44AHJDORF028	-3	9071..9235	54	
33	44AHJDORF030	3	14487..14648	53	
34	44AHJDORF031	2	11039..11191	50	
35	44AHJDORF135	3	693..842	49	
36	44AHJDORF033	-1	3646..3795	49	
37	44AHJDORF032	-2	9306..9455	49	
38	44AHJDORF034	-3	14000..14146	48	
39	44AHJDORF035	-3	13811..13957	48	
40	44AHJDORF036	-3	10019..10165	48	
41	44AHJDORF022	-3	8468..8611	47	
42	44AHJDORF037	1	14788..14931	47	
43	44AHJDORF038	-2	3528..3671	47	
44	44AHJDORF039	3	1743..1883	46	
45	44AHJDORF040	2	9740..9877	45	
46	44AHJDORF041	2	15836..15973	45	
47	44AHJDORF042	-1	5014..5151	45	
48	44AHJDORF043	-1	4402..4539	45	
49	44AHJDORF044	-2	12783..12917	44	
50	44AHJDORF149	-2	639..770	43	
51	44AHJDORF046	1	4891..5019	42	
52	44AHJDORF047	1	11911..12039	42	
53	44AHJDORF045	2	10655..10783	42	
54	44AHJDORF048	-3	15212..15340	42	
55	44AHJDORF049	3	5784..5909	41	
56	44AHJDORF050	3	13158..13283	41	
57	44AHJDORF051	-2	10944..11066	40	
58	44AHJDORF052	-3	14216..14338	40	
59	44AHJDORF053	3	3348..3467	39	
60	44AHJDORF054	3	7551..7670	39	
61	44AHJDORF055	3	15705..15821	38	
62	44AHJDORF056	1	5512..5625	37	
63	44AHJDORF057	2	10121..10231	36	
64	44AHJDORF058	3	10767..10877	36	

65	44AHJDORF164	-1	592..702	36	
66	44AHJDORF059	-2	8250..8360	36	
67	44AHJDORF060	-2	6147..6257	36	
68	44AHJDORF061	2	15551..15658	35	
69	44AHJDORF062	1	4285..4389	34	
70	44AHJDORF063	-3	9383..9487	34	
71	44AHJDORF065	1	5029..5130	33	
72	44AHJDORF064	2	2609..2710	33	
73	44AHJDORF066	-2	10380..10481	33	

Table 18

Predicted amino acid sequences

44AHJDORF001

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12627 atgggattactagaatgcgatcaatatacaaaacatgaacgtcgaatgattttatactgggatatagaacattagcgtacaat
1 M G L L E C M Q Y H K H E R R M I L Y W D I E T L A Y N
12543 aaagttaacggacgaaaaaaacaaacaaatataaaaacgttacttattctgtagcaattgggttggttaaatgggttatgaaatt
29 K V N G R K K P T K Y K N V T Y S V A I G W F N G Y E I
12459 gatgttgaagtatttccgagtttcgaatctttttatgcagcattttatagctatgtgaaaagacgtgatacaatcacaaaatca
57 D V E V P P S F E S F Y D A F Y T Y V K R R D T I T K S
12375 aaaacagatattatcatgattgcacataactgtaataaatacagataatcattttttacttaagacaccatcggttattttgat
85 K T D I I M I A H N C N K Y D N H F L L K D T M R Y F D
12291 aatattacacgcgaaaaatatatttttaaaatctgcagaaagaaatgaacacacattaaaaatgaaagaggctactattttagcc
113 N I T R E N I Y L K S A E E N E H T L K M K E A T I L A
12207 aaaaatcaaaatgtaatttttagaaaaacgtgttaaatcttcaatcaatttagatttaacaatgttttttaaatgggttttaaat
141 K N Q N V I L E K R V K S S I N L D L T M F L N G F K F
12123 aatattattgataactttatgaaaacaaacacacatttaggtaagaattacttgatgggttatttaaacagaa
169 N I I D N F M K T N T S I A T L G K K L L D G G Y L T E
12039 tcacaaacttaaaacagatttttaattatagcatttttgataaagataatgatagatgaatgatagtgagcctatgactatgctgtg
197 S Q L K T D F N Y T I F D K D N D M N D S E A Y D Y A V
11955 aaatgtttttgcaaaactcacacctgaacaacttacatacattcataatgacgtgattatattaggtatgtgcatattcattat
225 K C F A K L T P E Q L T Y I H N D V I I L G M C H I H Y
11871 agtgataatttccaaattttgactatacaaaatcaacattttcattgaaattatggaaattcttacttgcaataatgaaatgaca
253 S D I F P N F D Y N K L T F S L N I M E S Y L N N E M T
11787 cgttttcagttactcaaccaatatacaagattataaaatattcttatacacattatcatttccatgatatgaatttttatgactat
281 R F Q L L N Q Y Q D I K I S Y T H Y H P H D M N F Y D Y
11703 attaaatcattctatcgtgggtgttttaaatatgtataacaccaaatacataaacaactaattgatgagccttgtttttctatt
309 I K S F Y R G G L N M Y N T K Y I N K L I D E P C F S I
11619 gacatcaattcagagttatcctttatgtgtatcatgataaaatccaacatgggttatacttttcaagaaacttccagaaacca
337 D I N S S Y P Y V M Y H E K I P T W L Y F Y E H Y S E P
11535 acgtaataccttacttttttagatgatgacaattatttttattatataagattgataaagatgtatttaacgatdtattatta
365 T L I P T F L D D D N Y F S L Y K I D K D V F N D G D L L
11451 attaaaatataatcagctgtattacgtcaaatgattgtgaaataactataataatgataatgattacgttaataatcaatacaaat
393 I K I K S R V L R Q M I V K Y Y N N D N D Y V N I N T N
11367 acattaagaatgattcaagacattacgggtattgtgcatgacatatacgtgttaattcgtttgttatatgaatgtgaatac
421 T L R M I Q D I T G I D C M H I R V N S F V I Y E C E Y
11283 tttcatgcagtgatattatttttcaaaactattttatataaaacacaaaggtaagttaaaaaacaaatcaaatgacatcacct
449 F H A R D I I F Q N Y F I K T Q G K L K N K I N M T S P
11199 tacgactatcacattactgatgatatacaacgaacacccataactcaaatgaggaggttatgttatctaaagtctgttttaaatgga
477 Y D Y H I T D D I N E H P Y S N E E V M L S K V V L N G
11115 ttatatggcatacctgcattacgtttcacatttttaacttattccggttagatgataacaatgaactatacaaatcattaacggt
505 L Y G I P A L R S H F N L F R L D D N N E L Y N I I N G
11031 tacaacaaactgaacgtaataatatttctcattttgtcacatcacgtttcattgtataacttattggttctcttccaaatc
533 Y K N T E R N I L F S T F V T S R S L Y N L L V P F Q Y
10947 ttaacggaagtgaattgacgacaattttatttattgcgatactgatagtttgtatagaaatccggttgtaaacccttattg
561 L T E S E I D D N F I Y C D T D S L Y M K S V L V L N G
10863 aaccccgattttattcgaccgatagccttaggtaaatgggatattgaaaacgaacagatagataagatgtttgtactgaatcat
589 N P S L F D P I A L G K W D I E N E Q I D K M F V L N H
10779 aagaataatgcataatgaatggaatgaaagattaaaattgtcttctgctggtataccgaaaaacgctttatgatacaagcgtcgat
617 K K Y A Y E V N G K I K I A S A G I P K N A F D T S V D
10695 tttgaaacctttgtacgtgaacaattctttgacgggtgccattattgaaaacaataaaagtatctataatgagcaaggtacaata
645 F E T F V R E Q P F D G A I I E N N K S I Y N E G T I
10611 tcgatataatccgtctaaaactgaaattgtatgtggtaatgtatatgatgaatattttactgatgaacttaataatgaaacgtgaa
673 S I Y P S K T E I V C G N V Y D E Y F T D E L N M K R E
10527 tttatataaaagacgctagagaaaaatttcgaccatagtcattttgatgatattctttatattgaaagtacatcggttcattt
701 F I L K D A R E N F D H S Q F D D I L Y I E S D I G S F
10443 tcacttaacgacttatttccagttgaacgttcagtaacatacaaaatctgatttgcataatataaaacgtgaacatgatgaata
729 S L N D L F P V E R S V H N K S D L H I L K R E H D E I
10359 aaaaaaggaactgttaa 10342
757 K K G N C *
44AHJDORF002
3789 atggcatataatgaaaacgatttttaaatatttttgacatttcgtccatttttagacgaaattttataaaacgagagaacgttat
1 M A Y N E N D F K Y F D D I R P F L D E I Y K R E Y
3873 acacggtttttagcatgatagagcagattataataactaattcaaaatcatattatgattatatttcaagattatcaaaactaaatt
29 T P F Y D D R A D Y N T N S K S Y Y D Y I S R L S K I D I
3957 gatgtattagcacgtcgtattttgggactatgacaatgaattaaaaaacgtttcaaaaattgggacgacttaataatgaaagcattt
57 E V L A R R I W D Y D N E L K K R F K N W D D L M K A F
4041 ccagagcaagcgaagacttatttagaggttggttaaacgacggtacgattgacagttatttcatgacgaggttataaaatatt
85 C E A K K D L F R G W L N D G T I D S I I H D E F K K Y
4125 agcgcaggattaacatcggcattttgtcttatttaaggttactgaaatgaaacaaatgaatgacttttaaatcagaagttaaagac
113 S A G L T S A F A L F K V T E M K Q M N D F K S E V K D
4209 ttaataaagatattgaccgttttccgttaattgggtttgaattaaatgaagcgttgaaacaaagtttgatgggtttgtgtgtatt

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275

803 atggcacaacaatctacaaaaaatgaaactgcacttttagtagcaaaagtcagctaaatcagcgttacaaagattttaatcatgat
1 M A Q Q S T K N E T A L L V A K S A K S A L Q D F N H D
887 tattcaaaatcttggacatttggcgacaaatgggataattcaaatacaatgttcgaaacatttgtaataataattttatccct
29 Y S K S W T F G D K W D N S N T M F E T F V N K Y L F P
971 aagattaatgagactttattaatcgatattgcattaggttaactcggttaattggttagctaaagagcaagattttatggacaa
57 K I N E T L L I D I A L G N R F N W L A K E Q D F I G Q
1055 tatagtgaagaatcgtgattatggacacagtaaccaattcaatggacttatctaaaaatgaggaatttaattgtaaaactgaat
85 Y S E E Y V I M D T V P I N M D L S K N E E L M L K R N
1139 tatcccgatggcaactaagttatatgttaacgggaattgtgaagaaacaaaaattcacattaaacaacaatgatacagctttc
113 Y P R M A T K L Y G N G I V K K Q K F T L N N N D T R F
1223 aatttccaaacatttagcagcgcgaactaattacgcttttaggtgtatacaaaaagaaaaatttctgataattaatgattagaagaa
141 N F Q T L A D A T N Y A L G V Y K K K I S D I N V L E E
1307 aaagaaatcgctgcaatgttagttgattactcattgaatcaattatccgaaacaaatgtacgtaagcaacatcaaaagaagat
169 K E M R A M L V D Y S L N Q L S E T N V R K A T S K E D
1391 ttgacagcaaaagtttttgaagcaatcctaaacttacaacacacagtgctaaatataatgaagacatcgtgcatcaggtggt
197 L A S K V F E A I L N L Q N N S A K Y N E V H R A S G G
1475 gcaattggacaatatcaactgtatcaaaatgaagattgtgatttttaacaacagattcattaaaaatcttatcttttagat
225 A I G Q Y T T V S K L K D I V I L T T D S L K S Y L L D
1559 actaagattgcaaacacattccagattgcaggcattgattttcacagatcacgtttattagttttgacgacttaggtggcggtgtt
253 T K I A N T F Q I A G I D F T D H V I S F D D L G V P
1643 aaagtaacaaaagaatttaagttacaaaaccaagattcaattgactttttacgtgcgtatggagattatcaatcacaaattagga
281 K V T K E F K L Q N Q D S I D F L R A Y G D Y Q S Q L G
1727 gatacaattccagttggtgctgtatttacttatgtatctaaacttaagagtttactggcaacggtgaagaaatttaaacca
309 D T I P V G A V F T Y D V S K L K E F T G N V E E I K P
1811 aaatcagatttatatgcgtttatttttgatattatcaattaaatataaacggttacacaaaagggtatgttaaaaccaccattc
337 K S D L Y A F I L D I N S I K Y K R Y T K G M L K P P F
1895 cataacctgaatttgatgaagttacacactggattcattactattcatttaagccatttagtccatttttaataaaatttta
365 H N P E F D E V T H W I H Y Y S F K A I S P F F N K I L
1979 attactgaccaagatgtaaatccaaaccagaggaagaattacaagaataa 2029
393 I T D Q D V N P K P E E E L Q E *
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1 M N N D K R G L N V E L S K E I S K R V V E H R N R F K
2128 cgtcttatgttttaactcgtttatggaaattttaccgctactaatcaactataccaatcgtgatacgggttggtatagattttatt
29 R L M F N R Y L E F L P L L I N Y T N R D T V G I D F I
2212 cagttagaactcagctttaagacaaaaacattatgtatgtgtggtgaagctagaataaagcaaatatgatcttctggttatgta
57 Q L E S A L R Q N I N V V V G E A R N K Q I M I L G Y V
2296 aataacacttacttttaactcaagcaccaaaatttttcatcaaaactttaatttccaaatttcaaaaacgattaactaaagaagata
85 N N T Y F N Q A P N F S S N F N F Q F Q K R L T K E D I
2380 tattttatgtactgactatttaatacctgattgcttcaaaattcataagctatatgataactgtatgagtggttaacttt
113 Y P I V P D Y L I P D D C L Q I H K L Y D N C M S G N F
2464 gttgctcatgcaaaaataaacaattcaatataatgdtatagaataatagaacattatactgatgaattagcagaagtgct
141 V V M Q N K P I Q Y N S D I E I I E H Y T D E L A E V A
2548 ttatctcgctttttttaaactcatgcaagcaaaatttagcaagatatttaaatcagaaattaatgacgagtcattcaatcaactt
169 L S R F S L I M Q A K F S K I F K S E I N D E S I N Q L
2632 gtgtcggaaatataaacggtgcaccatttggtaaaatgtccacattgttttaatgcagatgacgatatttaaacaggt
197 V S E I Y N G A P F V K M S P M F N A D D D I I D L T S
2716 aatagcgttaatccagcatttaactgaaatgaaacgggaatatacaaaaacaaattagtgaaatgaagtaactatttaggcattaat
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2800 tcattagcgttgataaagaaagcgtgtttcagacgaagaggcaaaaagtaactcgtggatttaccacatcaaacagtaataatc
253 S L A V D K E S G V S D E E A K S N R G F T T S N S N I
2884 tattttaaagggtcgtgaaccaattacgtttttatcaaaagcgttatggttttagatattaaacggtattacgatgatgaacaacg
281 Y L K G R E P I T F L S K R Y G L D I K P Y Y D D E T T
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309 S K I S M V D T L F K D E S S D I N G *
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3104 aaattaacgttttatgatgatgaatttcaattcatgcaaaaaatgctgaagttcgacaaagacggttttagctatcgttaattgaa
29 K L T F Y D D E F Q F M Q K M L K F D K D V L A I V N E
3188 aaagatttttaagggtttttcattgaaagatgaattatcagatttacttttttaaaaaatcatttacgattcatttttagataga
57 K V F K G F S L K D E L S D L L F K K S F T I H F L D R
3272 gaaatcaacagacaaaacagttgaagcatttggcatgcaagtgattactgtatgtattacacatgaggattatttaaatggtggt
85 E I N R Q T V E A F G M Q V I T V C I T H E D Y L N V V
3356 tattcatcaagtgaagttgaaaaatcattacaatcacaagggttcacagaacacaatgaagatacaacaagtaacactgatgaa
113 Y S S E V E K Y L Q S G F T E H N E D T T S N Y L D E
3440 acatcgaatcaaaatgctacatctttagacaattcaactggcattgactgcaaacagaaacgcttatgtgtcattaccacaaagt
141 T S N Q N A T S L D N S T G M T A N R N A Y V S L P Q S
3524 gaggttaacattgatgttgataatacaacggttactcgtgataataacgattgataacggttaaaactgtgaataaatcg.
169 E V N I D V D N T T L R F A D N N T I D N G K T V N L K S
3608 agtaacgaaagtaatacaaacgcaaacgtaatacaaaatgaaaggttaacgaaaggtacacaattcactaagcagatttta
197 S N E S N Q N A K R N Q N Q K G N A K G T Q F T K Q Y L
3692 attgataatattgataaagcgtacgatttaagaaagaaaatttttaaatgaatttgataaaaaatgttttttacaatttggtag
3775
225 I D N I D K A Y D L R K K I L N E F D K K C F L Q I W *
44AHJDORF009

5744 atgaaatcacacaacaagcaaaagaatggatatataagcatgagggggcagggtgttgactttgatgggtgcatatggatttcaa
1 M K S Q Q Q A K E W I Y K H E G A G V D F D G A Y G P Q
5828 tgtatggacttatcagttgcttattgtgtattacattactgacggtaagttcgcattgtggggttaagctaaagacgcgataaat
29 C M D L S V A Y V Y Y I T D G K V R M W G N A K D A I N
5912 aatgactttaaagggttagcgacgggtgtataaaaatacaccgagctttaaacctcaattaggggacgtgtgctgtatatacaaat.
57 N D F K G L A T V Y K N T P S F K P Q L G D V A V Y T N
5996 ggacaatatggacatatccaatgtgtgttaagtggaaatcttgattattatacatgcttagaacaaaactgggttaggcggcggt
85 G Q Y G H I Q C V L S G N L D Y Y T C L E Q N W L G G G
6080 tttgacgggtgggaaaaagcaaccatttagaacacattattatgacgggtgaactcactttattagacctaaattttcaggtagt
113 F D G W E K A T I R T H Y Y D G V T H F I R P K F S G S
6164 aatagcaaacgatttagaacatcaaaagtaaatcatttggaaaaatggaaacgaaaccaatcggcacatatattatagaatgaa
141 N S K A L E T S K V N T F G K W K R N Q Y G T Y Y R N E
6248 aatggtagacattttagctgtgtttttacaaatatttgcacgtgtcggtagtccaaaattatcagaacctaatggctatttggttc
169 N G T F T C G F L P I F A R V G S P K L S E P N G Y W F
6332 caaccaaacggtatatacaccataatacgaagttgtttatcagatggttagctatggattgggtataactggcaaggcacacgt
197 Q P N G Y T P Y N E V C L S D G Y V W I G Y N W Q G T R
6416 tattattaccagtcgccaatggaaatggaaaaacaggttaattagttacagtggttggtattccttgggggtgttctcgtata 6496
225 Y Y L P V R Q W N G K T G N S Y S V G I P W G V F S *

44AHJDORF010
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14336 gatttttagtaattgtaatttttaattttgatgataaagatttacaagagcggtacattgacacatggaaacattttgacacatctg
29 D F S N V N F K F D D K D L Q E A Y I D T W K H F A H F
14252 ccctattttctaaagaaagaaacgtatcatatgtaaatgctgtatcatttgtaagaggttcaagacataaaaaatttaattat
57 P Y F P K E R N V S Y V N A V S L V R G S R H K K L N Y
14168 attcttgaaatatataacgtaattgatgatttctaataataaaaacgctaaaaagcataaatacgttttataattttacaagct
85 I L E I Y N R N D D S N N K N A K K H K Y A L Y N L Q A
14084 aaaaataaattcttcaatgtataaatatattaaagaaatcgatactttatataaagaaattggtaaatcagatagaccagtg
113 K N N N S S M Y K Y I K E I D T L Y K E I G K S D R P V
14000 acaaatattgatgatgaagatgtgaggtataactttttatattatgcaacatttgacgaataa 13938
141 T N I D D E D V R Y N F L Y Y A T F D E *

44AHJDORF011
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1 M T N V K D I L S R H Q N T T L A R F E F E E K E R E F I
15509 aaactatcagaatttagtagaaaaatacgggtatgaaaaagagtatatcgtagagcattattcacacaaagaaatcaaaattc
29 K L S E E L V E K Y G M K K E Y I V R A L F T N K E S K F
15425 ggtgaacaagggttctcgtcactgatgactataacgtaaaacttaccgaaccacttaacagaaatttaataaagaatgagagca
57 G E Q G V I V T D D Y N V N L P N H L T E L I K E M R A
15341 gatgaggacggttggtagattatcaatgtggagaagttcaattcacaattttatgaatatgaaaacaaaaaggtcaaaaaggt
85 D E D V V D I I N A G E V Q F T I Y E Y E N K K G Q K G
15257 tactcaatcaattttggtcaagtatcattttaa 15225
113 Y S I N F G Q V S F *

44AHJDORF012
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1 M N E V K F R F T D S E A P H M F I Y A G D L K L L Y F
8475 ttatttgtattaatgttctgttattatttacaggtattttcaaaagcaattaaaaataaacttatgggtcaaaaaatcaatg
29 L F V L M F V D I I T G I S K A I K N N N L W S K K S M
8559 agaggattttctaaaaaattattgatattctgtattatttagcaaacatttgaccagattttacaattaaaaggtggt
57 R G F S K K L L I F C I I I L A N I I D Q I L Q L K G G
8643 ctactcatgattacaatattttattatattgcaaatgagggtctttctattgtagaaaattgtgcagaatggacgtatttagta
85 L L M I T I F Y Y I A N E G L S I V E N C A E M D V L V
8727 ccagaacaaattaaagataaattaaagagtcattaaaaatgatactgaaaagagtgataacaaatgaacgatcaagagaagataga
113 P E Q I K D K L R V I K N D T E K S D N N E R S R E D R
8811 taa 8813
141 *

44AHJDORF013
14996 atgaaaaataaaactacttttagattaaataattttattaccttttaacaaatagagattattataatgataaatttgaa
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14912 aaatttacttcatctataaaaaatgtatagtaaaaaataatattgggtgatgtgtatattgagtttgacaacaaatattgatg
29 K F T S S N K K C I V K I N M G D V Y I E F D K Q Y D D
14828 tttgaaattgaaaaagagttatttacggttagatatacgacattgataaaaaaacatgttttaataacttgattttattat
57 F E I E K E L F T L D I D I D I K K H V F N I L V F Y Y
14744 agaaattatttaagtaattgaattaaagagaattttattaaacggttacaattgacgacgtattattcaaaattttgataaacct
85 R N Y L S N E L I R E I L L N V T I D D V L S N F D K P
14660 cttgaaagcgaattaatgatttttatcaaaaacaaagtcataacgataatgggaaagtgattgacatgaataa 14586
113 L E S E L M I I Y Q N K V I Y D N G K V I D H E *

44AHJDORF113
199 atgacagaattttagtgaatcgtaaaacagacgacaaagaagaacttcagaatcaactgaagaaaatttagaatacaactgaa
1 M T E F D E I V K P D D K E E T S E S T E E N L E S T E
283 gaaacttcagaatcaactgaagaatcaactgaagaatcaactgaagataaaacagtagaacaatcgtagaa
29 E T S E S T E S T E S T E S T E S T E D K T V E T I E E
367 gaaaaatgaaacaaatttagaacctactacaacagatgaagatagttcgaaatttgacctgtgtattagaacacgtattgct
57 E N E N K L E P T T T D E D S K F D P V V L E Q R I A
451 tcattagaaacaacagtgactctttttatcttcacaaatgcaacaaccacaacagtagaacaacacacacagatcagatgaaca
85 S L E Q Q V T T F L S S Q M Q Q Q Q V Q Q T Q S D V T
535 gaatacaacaaagaagataacgactattcagatgaagaactagttgataagtttagatttagattag 600

277

113 E S N K E D N D Y S D E E L V D K L D L D *
44AHJDORF114
16172 atgggttaattgtgataatgcaccagaagaaaaaggacaagcctatactgaaatgttgcaactattcaataaactgattcaatgg
1 M V N V D N A P E E K G Q A Y T E M L Q L F N K L I Q W
16088 aattccagcttatacatttgacaatgcaatttaacttattatcggttgccaacaactattattaaactataatagttctgtgtt
29 N P A Y T F D N A I N L L S A C Q Q L L L N Y N S S V V
16004 caattcttaaatgatgaactaaacaacgaaactaaaccagaatcaatattgtcttatattgctgtgtgacccaatagaacaa
57 Q F L N D E L N N E T K P E S I L S Y I A G D D P I E Q
15920 tgggaatgcatgataaaggattttatgaaacgtataacgtttacgttttttag 15870
85 W N M H K G F Y E T Y N V Y V F *
44AHJDORF014
6243 atgaaaatggtacattttacatgtggttttttaccatattttgcacgtgtcggttagtccaaaattatcagaacctaatggctatt
1 M K M V H L H V V F Y Q Y L H V S V V Q N Y Q N L M A I
6327 gggtcccaacaaacggttatcacccatataacgaagttgtttatcagatgggtacgtatggattgggtttggtgcaaggca
29 G S N Q T V I H H I T K F V Y Q M V T Y G L V I T G K A
6411 cacgttattatttaccagtgcgcaatggaatggaaaaacaggttaattagttacagtgttggtattccttggggggtgttctcat
57 H V I I Y Q C A N G M E K Q V I V T V L V F L G G C S H
6495 aatgggtatttttagcctttttctttga 6521
85 N G Y F S L F L *
44AHJDORF015
15403 gtgacgataaacacctgttccacgaatttttgattctttgtttgtgaataatgctctaacgatatactcttttttcataccgtat
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15487 tttttcactaattctgatagtttgataaattctcttttcttttctcctcaaatcctcctaatgtgttttgggtgtcttgat
29 F S T N S D S L S I N S L S F S S N S N L A N V F W C L D
15571 aaaaatctctttttagccttttttatttctcttctttaaattttgtcttctgcaattgcatgtag 15645
57 K I S P T F V I L F L L L F K L F A F C N C D L *
44AHJDORF016
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1 M K V D D I V T L R V K G Y I L H Y L D D D N E Y I E E
15768 tttttaccacttcacgagtatcatttaacaaaacacaaagcaaaagaattattaccagacacatgtaaaactattgtccactaca
29 F L P L H E Y H L T K T Q A K E L L P D T C K L L S T T
15684 cgcacaacgaaaacaattcaagtttattacaatgatttactacaatcgcaattgcagaaagcaataa 15616
57 R T T K T I Q V Y Y N D L L Q I A I A E S K *
44AHJDORF017
10757 atggaagattaaaattgtctctgtgtgtataccgaaaacgcctttgatataagcgtcgattttgaaacctttgtacgtgaac
1 M E R L K L L L L V Y R K T P L I Q A S I L K P L Y V N
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29 N S L T V P L L K T I K V S I M S K V Q Y R Y I R L K L
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57 K L Y V V M Y M M N I L L M N L I *
44AHJDORF018
1098 atgttaattggtactgtgtccataatcacgtattcttctactatattgtccaataaaatcttctcttttagctaaccaattaaaa
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29 R L P N A I S I N K V S L I L G N K Y L F T N V S N I V
930 tttgaattatccattttgtcgccaaatgtccaagattttgaataa 886
57 F E L S H L S P N V Q D F E *
44AHJDORF019
9836 atgttacctggtttgtataagtattcttttttgaataaaggtacaccaattgcttttttatatttttctggttaactgtgcatat
1 M L P G L Y K Y S F L N K G T P I A F L Y F S G N C A Y
9752 gtccagttaccaccaatcacacgaccactttttccattgtgctgactgatttaccactaattggttttatggtctccgcatca
29 V Q L P P I T R P L F P F G L T D L P L I G L W S P S S
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57 S V G L E L L L P L S T *
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1 M E N E T K N I E L K H V F R F K N G S L C I A L F D R
16278 acagaaaatgaaatttcttttatgatgttgacattgatgaaattgaagatttaaatcataattctgttttacgcgtaatttca
29 T E N E I S F Y D V D I D E I E D L N H N S V L R V I S
16194 actttattaggaagtataataatggttaa 16165
57 T L L G S D N N G *
44AHJDORF020
13865 atgtctaaacgattttgttttaccatgtttttgtccttgaatagtttatgatgtcggtttacagtgttaaaatttattcgtcaa
1 M S K R F C F T M F L L L V I V Y D V V Y S V K F I R Q
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29 M L H N I K S Y T S H L H H Q Y L S L V Y L I Y Q F L Y
14033 ataaagtacgatttctttaa 14053
57 I K Y R F L *
44AHJDORF123
614 atgtataggggaaacaacatgcgttctatgatgggtacatcatatgaagattcaagattaaataaacgaacagaaattaaatgaa
1 M Y E G N N M R S M M G T S Y E D S R L N K R T E L N E
698 aacatgtcaattgatacaataaaagtgaagatagttatgggtacaaaattcattcatttcaaaaacatttaccaggtgac
29 N M S I D T N K S E D S Y G V Q I H S L S K Q S P T G D
782 gttgaggaggaataa 796
57 V E E E *

44AHJDORF021
5816 atgacacatcaaagtcaacacctgccccctcatgcttatatatccattcttttgettggtgtgatttcatttatatcactc
1 M H H Q S Q H L P P H A Y I S I L L L V V V I S F I S L
5732 ctatttttgatgttttgcacccaacatattcacgatgttttgtttccgcattaacattactgaagaattctttatattccga
29 L F L M F C Y P T I F T M F C F R I N I T E E F F I F R
5648 tatattagcctctaa 5634
57 Y I S L *

44AHJDORF022
8611 atgtttgtcaaaatgataatacagaatatcaataattttttagaaaatcctctcattgatttttttgaccataagttattattt
1 M F A K M I I Q N I N N F L E N P L I D F F D H K L L F
8527 ttaattgcttttgaatacctgtaataatatcaacgaacattaatacaataaaaaagtag 8468
29 L I A F E I P V I I S T N I N T N K K *

44AHJDORF023
6494 atgagaacaccccccaaggaataacacactgtaactattacctgtttttccattccattggcgactggtaaaataaacgtg
1 M R T P P K E Y Q H C N Y Y L F F H S I G A L V N N N V
6410 tgccttgccagttataacaaatccatcgttaaccatctgataaacaacttcgttatatggtgataaacgtttggttggaacc
29 C L A S Y N Q S I R N H L I N K L R Y M V Y N R L V G T
6326 aatagccattag 6315
57 N S H *

44AHJDORF024
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1 V S M Y A S C K S L S S N L K L T L L K S F K N K S F S
14359 tgcctttttctagcttctctttttttttccatctatccatttcagacgtatgtctaaccaatgttatcaacctccatataag
29 C S F L A S L S F F H L S I S D V C L T N V I N L H I K
14443 cataaataa 14451
57 H K *

44AHJDORF025
15175 atggaacgtaaaatacaaaaacggtattattatattgcatgagattaaaggacattttccacatcaaactcattgtttgaagat
1 M E R K Y K T V L L Y C D E I K G H F P H Q I S M F E D
15091 ttatatgacgctaaagttgtatattcatattatgaatataacctgttccactaaaaataacgcttatcatagaatacattaag
29 L Y D A K V V Y S Y Y E Y N L F T K K Y A Y I I E Y I K
15007 gagatataa 14999
57 E I *

44AHJDORF026
14593 atgaataaacctattataacatagccattgttttcccttttagcatttttaattacacttatcatacttatgacactgcatatacgc
1 M N N L L N I A I V F L L A F L I T L I I L M T L H I R
14509 gtgtcattttggtgtttttattcactacattgatttatattcttatattatcttttttaaggtttatttatgcttttatatgaggttga
14426
29 V S F G V L F T T L I I F Y I I F L M V I Y A L Y G G *

44AHJDORF027
12916 atgattgtctatatccctaatttttagtataaaattcatattgttttgtatatggtacaacgataatattttgcataaaagtagt
1 M I V I P N F S T K F I L F C I W Y N D N I C H K S S
13000 tacattatacatgactttaatatatttatccatcagttttgatataagaataacacgttttgattgatgtgtattcttaa
13080
29 Y I I H D F N I F I I S F D I E E I T V L I D V I S *

44AHJDORF029
15183 gtgttttaaatggaacgtaaaatacaaaaacggtattattatattgcatgagattaaaggacattttccacatcaaactcattgt
1 V F K W N V N T K R Y Y Y I A M R L K D I F H I K S Q C
15099 ttgaagatttatgacgctaaagttgtatattcatattatgaatataacctgttccactaaaaataacgcttatatcatag
15019
29 L K I Y M T L K L Y I H I M N I T C S L K N T R I S *

44AHJDORF028
9235 atggaatatatgacgctcaattgtacctgctttcatatttttgcaaaatctgcattaccttttcttgtacgtcttggta
1 M E Y M H V Q L Y L L S Y F L Q N L H Y L F F V R L V V
9151 caaagtggacgatgttacctgcgtcatacaagacggtgtccagcttgttttgattgtgataactaactttcttgctatga 9071
29 Q S G R C Y L R H T K T V V Q L V L I V I L T F L L *

44AHJDORF030
14487 gtgaataaaaacaccaaataacacgcttatatgcagtgatcataagtgataagtgtaattaaaaatgctaaaaggaaaacaatg
1 V N K T P N D T R I C S V I S M I S V I K N A K R K T M
14571 gctattgttaaataggttattcatggtcaatcactttccattatcgatatgactttgttttgataaataatcattaa 14648
29 A M F N R L F M V N H F P I I V Y D F V L I N N H *

44AHJDORF031
11039 atgatattgtatagttcattgttatcatctaaacggaataagttaaaatgtgaacgtaatgcaggtatgccatataatccattt
1 M I L Y S S L L S S K R N K L K C E R N A G M P Y N P F
11123 aaaacgactttagataacataacctcctcatttgagtatgggtgttcgttgatatcatcagtaattgtga 11191
29 K T T L D N I T S S F E Y G C S L I S S V M *

44AHJDORF135
693 atgaaaacatgtcaattgtatacaaaaagtgaaagatagttatggtgtacaaaattcattcactttcaaaacaattcattacag
1 M K T C Q L I Q I K V K I V M V Y K F I H F Q N N H L Q
777 gtgacgttgaggaggaataataaattatggcacaacaactacaaaaaatgaaactgcacttttag 842
29 V T L R R N N K L W H N N L Q K M K L H F *

44AHJDORF033
3795 atgccattatattaaccacctctaccaaaattgttaaaaacattttttatcaaattcatttaaaattttctttcttaaatcgta
1 M P L F N H L Y Q I C K K H F L S N S F K I F F L K S Y

3711 gctttatcaatattatcaattaaatactgcttagtgtaattgtgtaccttttgcattacctttttga 3646
29 A L S I L S I K Y C L V N C V P F A L P F *
44AHJDORF032
9455 atggctgttttgcataaagcgagtagtgtaactaccactgtcaccactactaccactgtcagacgaatcactaggtgatccacct
1 M A C F A K A S S E L P L S P L L P L S D E S L G D P P
9371 ttaccgtctaattaccaccccaagctagaatagtagtattcgaccgtctaaaaatggattaccatag 9306
29 L P S N L P P Q A R I V F A P S K N G L P *
44AHJDORF034
14146 atgatgattctaataataaaaacgctaaaaagcataaatacgtttatataatttacaagctaaaaataataattcttcaatgt
1 M M I L I I K T L K S I N T L Y I I Y K L K I I I L Q C
14062 ataaatatattaagaatcgatactttatataaagaattggtaaatcagatagaccagtga 14000
29 I N I L K K S I L Y I K K L V N Q I D Q *
44AHJDORF035
13957 atgcaacatttgacgaataaatttaacactgtaaacgacatcataaactattacaaggagcaaaaacatggtaaaaacaaatcg
1 M Q H L T N K F N T V N D I I N Y Y K E Q K H G K T K S
13873 ttttagacatggtaagagattatcaaaatgctgtcaatcatgtcagaaaaaaatcccagataa 13811
29 F R H G K R L S K C C Q S C Q K K N P R *
44AHJDORF036
10165 gtgtatacaataccacacgtgatgggtgcaacatattgggtgtacattatagtttgcaactaaaaacgaacctcttcaaaaactg
1 V Y T I P H V M V Q H M V V H Y S L Q L K T N H L Q K L
10081 ctacaacaacacgtgtgtgaccaataccatattgcagttgctgtgaagtattgggtgtttactag 10019
29 L Q Q H L C D Q Y H M Q L L V S M V V Y *
44AHJDORF037
14788 atgtcgatatctaacgtaataaactctttttcaatttcaaaatcatcatattgtttgtcaaaactcaatatacacatcacccata
1 M S I S N V N N S F S I S K S S Y C L S N S I Y T S P I
14872 tttatttttactatacattttttattagatgaagtaatttttcaattttatcattataa 14931
29 F I F T I H F L L D E V N F S N L S L *
44AHJDORF038
3671 gtgtaccttttgcattacctttttgattttgattacgttttgcgttttgcattactttcgttactcgatttttccaggttttac
1 V Y L L H Y L F D F D Y V L R F D Y F R Y S I Y S Q F Y
3587 cgttatcaatcgattattattatcagcgaatcgtaacgtgtgtattatcaacatcaatgttaa 3528
29 R Y Q S Y Y Y Q R I V T L Y Y Q H Q C *
44AHJDORF039
1743 gtgctgtatttacttatgatgtatctaaacttaagaggtttactggcaacgttgaagaaattaaacaaaatcagatttatatg
1 V L Y L L M M Y L N L K S L L A T L K K L N Q N Q I Y M
1827 cgtttttttggatattaattcaattaaataaaacgttacacaaaaggatgttaa 1883
29 R L F W I L I Q L N I N V T Q K V C *
44AHJDORF040
9740 gtggtaactggacatatgcacagttaccagaaaaataaaaaagcaattgggtgtaccttttattcaaaaaagaatacttatata
1 V V T G H M H S Y Q K N I K K Q L V Y L Y S K K N T Y T
9824 aaccaggtaacatatcttctcaaacgggttaatgcaggacaatgtacagaattaa 9877
29 N Q V T Y F L K R V M Q D N V Q N *
44AHJDORF041
15836 atgtcgtcaactttcattattatcatcactccttttcaaaaaacgttaaacgttatcgttttcataaaatcctttatgcattatcc
1 M S S T F I I I S L L S K K R Y T F H K I L Y A Y S
15920 attgttctattgggtcatcaccgaataataagacaattattgattctggttttag 15973
29 I V L L G H H Q Q Y K T I L I L V *
44AHJDORF042
5151 atgcacgacgcgtcgtcttttgttaatttatagttttgtgaacctcttgcgcgtaatgcttcaaagtgttcatactcaccaagtt
1 M H D R R L L L I Y S F V N L L R V M L Q S V H T H Q V
5067 ggaagaaccatataaattatggaaacgttttccaccacgcggtttgtcatag 5014
29 G R N H I N Y G N V F H H R R L S *
44AHJDORF043
4539 atgcgacttgtaacagttttgcaacaccatcgatgtaacagatttttcatttcaccattggattgacgttctaataccgattg
1 M R L V T V L Q H H R D V T R F S F H H W I D V L I R L
4455 ttgtaccatgaccaccctgtacaatcgcgtgcttgaattaagtcaccactag 4402
29 L Y H D H P V Q Y A C L K L S H H *
44AHJDORF044
12917 atgttacctatttactgtgatgatgttttataaagaaaacatggaacgttattactacaatccaagcaatttacattttgaca
1 M L P I Y V M I C F I K K T W N V I T T I Q A I Y I L T
12833 atgcttactctaaaaattacgtggttgataatgatagattttatatttag 12783
29 M L T L K I T W L I M I D I Y I *
44AHJDORF149
770 atgattgttttgaagtgaaatgaattgtacaccataactatcttcaacttttatttattgtatcaattgacatgttttcatttaatt
1 M I V L K V N E F V H H N Y L H F Y L Y Q L T C F H L I
686 ctgttcgtttattttaattcttgaatcttcatatgatgtacccatcatag 639
29 L F V Y L I L N L H M M Y P S *
44AHJDORF046
4891 atgattatccatttaagtattcatatcaagacggtattaatttcccacgtgataaactttaagagcctgaagggtatttgcattt
1 M I I H L S Y H I K T V L I S H V I T L K S L R V~F A F
4975 atacaatccaaaaacaaacgtaaatcggttattacttgcattatga 5019
29 I Q I Q K Q N V N R Y Y L L *
44AHJDORF047
11911 atgaatgtatgtaagttgttcagggtgtgagttttgcaaaacatttcacagcatagtcaggttccattatcattcatattcatt
1 M N V C K L F R C E F C K T F H S I V I G F T I I H I I

11995 atctttatcaaaaatcgataaataaatctgttttaagtgtga 12039
29 I F I K N R I I K I C F K L *
44AHJDORF045
10655 atggcaccgtcaaagaattgttcacgtacaaagggttcaaaatcgacgcttgatcaaaggcgttttcggtataccagcagaa
1 M A P S K N C S R T K V S K S T L V S K A F F G I P A E
10739 gcaattttaattctttccattcattcatatgcatatttcttatga 10783
29 A I L I F P F T S Y A Y F L *
44AHJDORF048
15340 atgaggacgtgttgacattatcaatgctggagaagttcaattcacaattatgaatatgaaaacaaaaagggtcaaaaagggt
1 M R T L L T L S M L E K F N S Q F M N M K T K K V K K V
15256 actcaatcaattttgggtcaagtatcattttaatacaatttcataag 15212
29 T Q S I L V K Y H F N T I S *
44AHJDORF049
5784 atgaggggggcagggtgttgactttgatgggtgcatatggatttcaatgtatggacttatcagttgcttatgtgtattacattactg
1 M R G Q V L T L M V H M D F N V W T Y Q L L M C I T L L
5868 acggtaaagtctgcatgtgggtgaatgctaaagacgcgataa 5909
29 T V K F A C G V M L K T R *
44AHJDORF050
13158 gtgtgttactgtttttcattcacgtaatcggttctgctgcatttctaaaaaaatgtttttgtaaagtccttgatgtattcattttat
1 V C Y V F H S R N R F V A F L K K C F C K V L M Y S F Y
13242 gcttttgtaataaattgtatatattttaattggataatatag 13283
29 A F V I N C I Y L N W I I *
44AHJDORF051
11066 atgataacaatgaactatacaatatcattaacgggttacaaaaacactgaacgtaatatattattctctacatttgcacatcac
1 M I T M N Y T I S L T V T K T L N V I Y Y S L H L S H H
10982 gtccattgtataacttattgttcttttccaatacttaa 10944
29 V H C I T Y W F L S N T *
44AHJDORF052
14338 atgatttttagtaattgttaatttttaatttgatgataaagatttacaagggcggtacattgacacatggaacattttgcacatc
1 M I L V M L I L N L M I K I Y K R R T L T H G N I L H I
14254 tgccctattttcctaagaaagaaacggtatcatatgtaa 14216
29 C P I F L K K E T Y H M *
44AHJDORF053
3348 atgtgggtttattcatcaagtgaagttgaaaaatacttacaatcacaaggcttcacagaacacaatgaagatacaacaagtaaca
1 M W F I H Q V K L K N T Y N H K A S Q N T M K I Q Q V T
3432 ctgatgaaacatcgaaatcaaatgctacatctttag 3467
29 L M K H R I K M L H L *
44AHJDORF054
7551 atgactggaatggaataacgatgttactcgacgctggttaagatttcacaaaaaactgggtgtaagttacgtacaaaatcaatta
1 M T G M E I R C Y S T L V R F H K K L V L S Y V Q N Q L
7635 ttggttatcataatgaagttcgagtatatccagtag 7670
29 L V I I M K F E Y I Q *
44AHJDORF055
15705 atgtgtctggtaataattcttttgcctgtgttttgggttaaatgatactcgtaagtggttaaaaatccctaatgtattcattat
1 M C L V I I L L L V F W L N D T R E V V K I P Q C I H Y
15789 catcatctaagtaagtaagtatataacctttga 15821
29 H H L S N E V Y N L *
44AHJDORF056
5512 gtgagtattacattacaggttaaccaaatggaattatttagagacgcgccagaagaaattaaaaagtggtgcatggttacgtg
1 V S I T L Q V T K W N Y L E T R Q K K L K K W V H G Y V
5596 tgcgaagtggtaacgcagtcggtgaagtaa 5625
29 C Q V V T Q S V K *
44AHJDORF057
10121 atgtaccaccatattgttgcaccatcacgtgtggtattgtatacactcattaatggcggtaccaaataatgctggtgataatattg
1 M Y H H M L H H H V W Y C I H S L M A Y Q I M L V I I L
10205 tattcttttagtggtattgcttaattaa 10231
29 Y S L V V L L N *
44AHJDORF058
10767 atgcatatttcttatgattcagttacaaacatcttatctatctgttctgttttcaatatccatttacctaagggtatcggtcgga
1 M H I S Y D S V Q T S Y L S V R F Q Y P I Y L R L S G R
10851 ataaactgggttcaataagggttttaa 10877
29 I N W G S I R V *
44AHJDORF164
702 atgttttcatttaattctgttctgtttattttaattcttgaattcttcatatgatgtaccatcatagaacgcgatgtgtttccctca
1 M F S F N S V R L P N L E S S Y D V P I I E R M L F P S
618 tacatgtttaaatctctcctaattctaa 592
29 Y M F K F L L I *
44AHJDORF059
8360 atggattttgttaacattggattacctgaacgctcattatgccaaaatcttacaccagattctaaaattgcttttaattgttcca
1 M D F V T L D Y L N R H Y A K I L H Q I L K L L I V P
8276 ttaacatgggtcgatgtcacgtatag 8250
29 L T W G R C H V *
44AHJDORF060
6257 atgtaccattttcattttctataatatgtgccgtattggtttctgtttccattttccaaatgtatttacttttgatgttttcaatg
1 M Y H F H F Y N M C R I G F V S I F Q M Y L L L M F L M

6173 ctttgctattactacctgaaaatttag 6147
29 L C Y Y Y L K I *
44AHJDORF061
15551 atgtgttttgggtgtcttgataaaaatcttttacgtttgtcattttatttctcctcttatttaaattatttgctttctgcaatt
1 M C F G V L I K Y L L R L S F Y F S S Y L N Y L L S A I
15635 gcgatttgtagtaaatcattgtaa 15658
29 A I C S K S L *
44AHJDORF062
4285 gtgggtattcgcaacgcagttaaccaatctattaatattgataaagaacaaatcacatgtactctacacaatccgattctcaaa
1 V V F A T Q L T N L L I L I K K Q I T C T L H N P I L K
4369 aacctgaagggttttgggataa 4389
29 N L K V F G *
44AHJDORF063
9487 atgcgtcttgtatttttttaataattcttgcatggcctgttttgctaaagcgagtagtgaactaccactgtcaccactactac
1 M R L V F F L I I L A W L V L L K R V V N Y H C H H Y Y
9403 cactgtcagacgaatcactag 9383
29 H C Q T N H *
44AHJDORF065
5029 gtgggtggaaaacgtttccataatttatatggtttcttccaacttggtgagtatgaacactttgaagcattacgcgcaagaggtt
1 V V E N V S I I Y M V S S N L V S M N T L K H Y A Q E V
5113 cacaaaactataaattaa 5130
29 H K T I N *
44AHJDORF064
2609 atgacgagtcaatcaatcaacttgtgtccgaaatatataacgggtgcaccatttgttaaaatgtcacctatgtttaatgcagatg
1 M T S Q S I N L C P K Y I T V H H L L K C H L C L M Q M
2693 acgatatcattgatttaa 2710
29 T I S L I *
44AHJDORF066
10481 atgatattctttatattgaaagtgcacatcggttcattttcacttaacgacttatttccagttgaacgttcagtacataacaaat
1 M I F F I L K V T S V H F H L T T Y F Q L N V Q Y I T N
10397 ctgatttgcataatattaa 10380
29 L I C I Y *

Table 19

Sequence similarities between ORFs 44AHJD and public databases

Phage: 44AHJD

Database: nr

Query= sid|110871|lan|44AHJDORF001 Phage 44AHJD ORF|10342-12627|-1
(761 letters)

gi 118848 sp P19894 DPOL_BPM2 DNA POLYMERASE >gi 76896 pir JQ0...	55	1e-06
gi 1072656 pir S51275 DNA polymerase - phage CP-1 >gi 836593 e...	53	6e-06
gi 1429230 emb CAA67649 (X99260) DNA polymerase [Bacteriophage...	49	1e-04
gi 1572479 emb CAA65712 (X96987) DNA polymerase [Bacteriophage...	46	0.001
gi 118851 sp P06950 DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP...	45	0.002
gi 2435429 (AF012250) unassigned reading frame (possible DNA po...	45	0.002
gi 1084487 pir S41618 DNA polymerase - slime mold (Physarum po...	45	0.002
gi 4877819 gb AAD31446.1 (AF133505) DNA polymerase [Neurospora...	44	0.004
gi 461962 sp P33537 DPOM_NEUCR PROBABLE DNA POLYMERASE >gi 2833...	44	0.004
gi 2499511 sp Q12471 6P22_YEAST 6-PHOSPHOFRUCTO-2-KINASE 2 (PHO...	41	0.041
gi 2258375 gb AAD11909.1 (AF007261) transcription initiation f...	40	0.070
gi 15734 emb CAA37450 (X53370) DNA polymerase (AA 1-575) [Bact...	39	0.092

Query= sid|110872|lan|44AHJDORF002 Phage 44AHJD ORF|3789-5732|3
(647 letters)

gi 135273 sp P27622 TAGC_BACSU TEICHOIC ACID BIOSYNTHESIS PROTE...	112	7e-24
gi 142847 (M64050) DNase inhibitor [Bacillus subtilis]	52	1e-05
gi 4038407 (AF103943) factor C protein precursor [Streptomyces ...	39	0.10

Query= sid|110873|lan|44AHJDORF003 Phage 44AHJD ORF|6626-8389|2
(587 letters)

gi 138123 sp P04331 VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9) >...	92	8e-18
gi 138124 sp P07534 VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) >...	82	1e-14
gi 1429238 emb CAA67657 (X99260) tail protein [Bacteriophage B...	78	2e-13
gi 215339 (M12456) p9 tail protein [Bacteriophage phi-29] >gi 2...	71	2e-11
gi 1181968 emb CAA87738.1 (Z47794) tail protein [Bacteriophage...	54	3e-06
gi 1181970 emb CAA87740.1 (Z47794) tail protein [Bacteriophage...	42	0.010

Query= sid|110875|lan|44AHJDORF005 Phage 44AHJD ORF|12643-13890|-1
(415 letters)

gi 3845203 (AE001399) GAP domain protein (cyclic nt signal tran...	52	6e-06
gi 3758843 emb CAB11128.1 (Z98551) predicted using hexExon; MA...	49	5e-05
gi 3845297 (AE001421) hypothetical protein [Plasmodium falciparum]	48	1e-04
gi 4493936 emb CAB38972.1 (AL034556) predicted using hexExon; ...	47	2e-04
gi 3845165 (AE001390) hypothetical protein [Plasmodium falciparum]	46	6e-04

Query= sid|110877|lan|44AHJDORF007 Phage 44AHJD ORF|2044-3027|1
(327 letters)

gi 1181960 emb CAA87731.1 (Z47794) connector protein [Bacterio...	46	5e-04
gi 1429239 emb CAA67658 (X99260) upper collar protein [Bacteri...	45	8e-04
gi 137915 sp P07535 VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR ...	44	0.002
gi 137914 sp P04332 VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ...	41	0.009

Query= sid|110878|lan|44AHJDORF008 Phage 44AHJD ORF|3020-3775|2
(251 letters)

gi 4982468 gb AAD30963.2 (AF118151) SNF1/AMP-activated kinase ...	52	3e-06
gi 1730077 sp P18160 KYK1_DICDI NON-RECEPTOR TYROSINE KINASE SP...	46	2e-04
gi 3758855 emb CAB11140.1 (Z98551) predicted using hexExon; MA...	46	2e-04
gi 585795 sp P21538 REB1_YEAST DNA-BINDING PROTEIN REB1 (QBP) >...	46	3e-04
gi 172372 (M58728) DNA-binding protein [Saccharomyces cerevisiae]	46	3e-04
gi 2952545 (AF051898) coronin binding protein [Dictyostelium di...	45	6e-04
gi 535260 emb CAA82996 (Z30339) STARP antigen [Plasmodium reic...	45	7e-04
gi 1429240 emb CAA67659 (X99260) lower collar protein [Bacteri...	44	0.001

Query= sid|110879|lan|44AHJDORF009 Phage 44AHJD ORF|5744-6496|2
(250 letters)

gi 2764981 emb CAA69021.1 (Y07739) N-acetylmuramoyl-L-alanine ...	180	1e-44
gi 113675 sp P24556 ALYS_STAAU AUTOLYSIN (N-ACETYLMURAMOYL-L-AL...	118	6e-26
gi 1763243 (U72397) amidase [bacteriophage 80 alpha]	118	6e-26
gi 4574237 gb AAD23962.1 AF106851_1 (AF106851) LytN [Staphyloco...	84	9e-16
gi 3767593 dbj BAA33856.1 (AB015195) LytN [Staphylococcus aureus]	84	9e-16
gi 2764983 emb CAA69022.1 (Y07740) cell wall hydrolase Ply187 ...	77	2e-13
gi 3287732 sp O05156 ALE1_STACP GLYCYL-GLYCINE ENDOPEPTIDASE AL...	73	2e-12
gi 79926 pir A25881 lysostaphin precursor - Staphylococcus sim...	69	3e-11
gi 126496 sp P10548 LSTP_STAST LYSOSTAPHIN PRECURSOR (GLYCYL-GL...	69	3e-11
gi 3287967 sp P10547 LSTP_STASI LYSOSTAPHIN PRECURSOR (GLYCYL-G...	69	3e-11
gi 3341932 dbj BAA31898.1 (AB009866) amidase [peptidoglycan hy...	68	6e-11

Query= sid|110882|lan|44AHJDORF012 Phage 44AHJD ORF|8391-8813|3
(140 letters)

gi 140528 sp P24811 YQXH_BACSU HYPOTHETICAL 15.7 KD PROTEIN IN ...	80	6e-15
gi 4126631 dbj BAA36651.1 (AB016282) ORF45 [bacteriophage phi-...	76	1e-13
gi 141088 sp P26835 YNGD_CLOPE HYPOTHETICAL 14.9 KD PROTEIN IN ...	61	4e-09
gi 2293160 (AF008220) YtkC [Bacillus subtilis] >gi 2635548 emb ...	36	0.099
gi 1181973 emb CAA87743.1 (Z47794) holin protein [Bacteriophag...	31	3.3

Table 20

Homologies between phage 44 AHJD ORFs and proteins in public databases

Query= pt|110871 44AHJDORF001 Phage 44AHJD ORF |10342-12627|-1 1
(761 letters)

>gi|118848|sp|P19894|DPOL_BPM2 DNA POLYMERASE >gi|76896|pir||JQ0161
DNA-directed DNA polymerase (EC 2.7.7.7) - phage M2
>gi|215509 (M33144) DNA polymerase [Bacteriophage M2]
Length = 572

Score = 55.4 bits (131), Expect = 1e-06
Identities = 96/426 (22%), Positives = 159/426 (36%), Gaps = 88/426 (20%)

Query: 229 KLTPEQLTYIHNDVIIILGMCHIHYSDIFPNFDYNKLTFSNLNIMESYLNEMTR-----FQ 283
++TPE+ YI ND+ I+ DI +++T + ++ + + T+ F
Sbjct: 154 EITPEEYIYKNDIEIARA----LDIQFKQGLDRMTAGSDSLKGFKDILSTKKFNKVFP 209

Query: 284 LLNQYQDIKISYTHYHFDNMNFYDIKSFYRGGLNMYNTKYINKLIDEPFCSIDINSSYP 343
L+ D +I + YRGG N KY K I E D+NS YP
Sbjct: 210 KLSLPMDEI-----RKAYRGGFTWLNKYKEKEIGEGMV-FDVNSLYP 252

Query: 344 YVMYHEKIPTWLYFYEYSEPTLIPTFLDDDNYSLSYKIDKDVFNDDLLIKIKSRVLRQM 403
MY +P Y P + + D + LY I + F +L K + +
Sbjct: 253 SQMYSRPLP-----YGAPIVFQGYEKDEQYPLY-IQRIRFEFEL----KEGYIPTI 299

Query: 404 XXXXXXXXXXXXXXXXXXXXLRMIQ-DITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
+ ++ +T +D I+ + + +Y EY F +
Sbjct: 300 QIKKNPFKNGEYLNKSGVEPVLYLTNVDELILQEH-YELYNVEYIDGFK-----FRE 352

Query: 463 TQGLKLNKINMTSPYDYHTDDINEHPYSNEEVMLSKVVLNGLYG-----IPAL 511
G K+ I+ + H + L+K++LN LYG +P L
Sbjct: 353 KTGFLFKDFIDKWTYVKTH-----EEGAKKQLAKMLNLSLYGKFASNPVDTGKVPYL 403

Query: 512 RSHFNL-FRLDDNNELNYNIINGYKNTERNILFSTFVTSRSLYNLLVPPQYLTSEIDDNF 570
+ +L FR+ D YK+ + F+T+ + + + Q D
Sbjct: 404 KDDGSLGFRVGDDE-----YKDPVYTPM-GVFITAWARFTTITAAQACY-----DRI 449

Query: 571 IYCDTDSLYMKSVVKPLNPSLFDPIALGKWDIENEQIDKMFVLNHHK-----YAYEVNG 625
IYCDTDS+++ P + + DP LG W E+ + L K Y EV+G
Sbjct: 450 IYCDTDSIHLTGTEVPEIHKDIVDPKGLGYWAHES-TPKRAKYLRQKTYIQDIYVKEVDG 508

Query: 626 KIKIAS 631
K+K S
Sbjct: 509 KLKECS 514

>gi|1072656|pir||S51275 DNA polymerase - phage CP-1
>gi|836593|emb|CAA87725.1| (247794) DNA polymerase
[Bacteriophage CP-1]
Length = 568

Score = 53.5 bits (126), Expect = 6e-06
Identities = 104/464 (22%), Positives = 169/464 (36%), Gaps = 66/464 (14%)

Query: 230 LTPEQLTYIHNDVIIIL--GMCHIHYSDIFPNFDYNKLTFSNLNIMESYLNEMTRFQLLNQ 287
+ PE + YIH DV IL G+ ++Y + F Y + +L + +F+
Sbjct: 152 IKPEWIDYIHVDVAILARGIFAMYEEENFTK--YTSASEALTEFKRIFRKSRRKFRDFFP 209

Query: 288 YQDIKISYTHYHFDNMNFYDIKSFYRGGLNMYNTKYINKLIDEPFCSIDINSSYPYVMY 347
D K+ D+ + G + K+ + +++ DINS YP M
Sbjct: 210 ILDEKVD-----DFCRKHIVGAGRLPTLKHGRGRTLNQLIDIYDINSYMPATML 257

Query: 348 HEKIPTWLYFYEYSEPTLIPTFLDDDNYSLSY-KIDKDVFNDDL-LIKIKSRVLRQMXX 405
+P + + Y P + +D+Y+ + K D D+ L I+K ++
Sbjct: 258 QNALPIGIP--KRYGK---PKEIKEDHYIYHIKADFDLKRGLPTIQIKKLDALRIG 312

Query: 406 XXXXXXXXXXXXXXXXXXXXLRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 465
L + + H + E F +F +Y
Sbjct: 313 VRTSDYVTTSKNEVIDLYLTNFDLFLKHYDATIMYVETLE-FQTSDFDDYI----- 366

285

Query: 466 KLKKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYGPALR--SHFNLFRLDDN 523
 + Y Y E+ S E +K++LN LYG + S L LDD
 Sbjct: 367 -----TTYRYK-----KENAQSPAQKQAKIMLSLYGKFGAKIISVKKLAYLDDK 412

Query: 524 NELNYIINGYKNTERNIL-----FSTFVTSRSLYNLLVFPQYLTESEIDNFIYCDTDS 577
 L +KN + + FVTS + + + Q E DNF+Y DTDS
 Sbjct: 413 GILR-----FKNDDEEEVQPVYAPVALFVTSIARHFIISNAQ-----ENYDNFLYADTDS 462

Query: 578 LYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHHKYYAYEVNGKIKIASAGIPKN 637
 L++ +L+ DP GKW E + K L K Y E+ + + K
 Sbjct: 463 LHLFHSDSLVD---IDPSEFGKWAHEGRAV-KAKYLRSLKYIEELIQEDGTTDLVD-KG 517

Query: 638 AFDTSVDFETFVREQFFDGAIENNKSIYNEQGTISIYPSKTEI 681
 A T E E F GA E ++ +G IY + +I
 Sbjct: 518 AGMTPEIKEKITFENFVIGATPEGKRASKQIKGGTLIYETTFKI 561

>gi|1429230|emb|CAA67649| (X99260) DNA polymerase [Bacteriophage
 B103]
 Length = 572

Score = 49.2 bits (115), Expect = 1e-04
 Identities = 93/422 (22%), Positives = 155/422 (36%), Gaps = 88/422 (20%)

Query: 229 KLTPEQLTYIHNDVIIILGMCHIHYSIDIPNFDYNKLTFSNLIMESYLNEMTR-----FQ 283
 ++TPE+ YI ND+ I+ DI +++T + + + + T+ F
 Sbjct: 154 EITPEEYKIKNDIEIARA-----LDIQFKQGLDRMTAGSDSLKGFKDILSTKKFNKVPF 209

Query: 284 LLNQYQDIKISYTHYFHDNMNFYDIKSFYRGGLNMYNTKYINKLIDPCFSDINSSYP 343
 L+ D +I + YRGG N KY K I E D+NS YP
 Sbjct: 210 KLSLPMDEI-----RRAYRGGFTWLNDKYKEKEIGGMV-FDVNSLYP 252

Query: 344 YVMYHEKIPTWLYFYEHYSEPTLIPTFLDDNYFSLYKIDKDVFNDDLLIKISRVLRQM 403
 MY +P Y P + + D + LY I + F +L K + +
 Sbjct: 253 SQMYSRPLP-----YGAPIVFQGYEKDEQYPLY-IQRIRPEFEL----KEGYIPTI 299

Query: 404 XXXXXXXXXXXXXXXXXXXXLRMIQ-DITGIDCMHIRVNSFVIYECYFPHARDIIFQNYFIK 462
 ++ +T +D I+ + + +Y EY F +
 Sbjct: 300 QIKKNPFPGNEYLKNSGAEPVELYLTNVDELIEQH-YEMYNVEYIDGPK-----FRE 352

Query: 463 TQGLKKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLY-----IPAL 511
 G K I+ + H + L+K++ + LYG +P L
 Sbjct: 353 KTGLFKFIDKWTYVYKTH-----EKGAKQLAKLMFDSLYGKFASNPDVTKVPYL 403

Query: 512 RSHFNL-FRLDDNNELNYIINGYKNTERNILFSTFVTSRSLYNLLVFPQYLTESEIDNFI 570
 + +L FR+ D YK+ + F+T+ + + + Q D
 Sbjct: 404 KEDGSLGFRVGDDE-----YKDPVYTPM-GVFTITAWARFTTITAAQACY-----DRI 449

Query: 571 IYCDTDSLYMKSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHHK-----YAYEVNG 625
 IYCDTDS+++ P + + DP LG W E+ + L K YA EV+G
 Sbjct: 450 IYCDTDSIHLTGTEVPEIHKDIVDPKLGWYAHES-TFKRAKYLKQKTYIQDIYAKEVDG 508

Query: 626 KI 627
 K+
 Sbjct: 509 KL 510

>gi|1572479|emb|CAA65712| (X96987) DNA polymerase [Bacteriophage
 GA-1]
 Length = 578

Score = 46.1 bits (107), Expect = 0.001
 Identities = 80/376 (21%), Positives = 146/376 (38%), Gaps = 54/376 (14%)

Query: 234 QLTYIHNDVIIILGMCHIHYSIDIPNFDYNKLTFSNLIMESYLNEMTRFQLLNQYQDIKI 293
 ++ Y+ +D++I+ + +F N D+ +T + + +Y EM + +Y +
 Sbjct: 162 EIEYLKHDLIIVALA---LRSMFDN-DFTSMTVGSDALNTY--KEMLGKQWKEKYFPVL- 214

Query: 294 SYTHYFHDNMNFYDIKSFYRGGLNMYNTKYINKLIDPCFSDINSSYPVYHEKIPT 353
 + I+ Y+GG N KY + + D+NS YP +M ++ +P
 Sbjct: 215 -----SLKVNSEIRKAYKGGFTWVNPYQGETVYGGMV-FDVNSMYPAMMKNKLLP- 264

Query: 354 WLYFYEHYSEPTLIPTFLDDNYFSLYKIDKDVFNDDLLIKISRVLRQMXXXXXXXXXX 413
 Y EP + + + LY F + KI ++

286

Sbjct: 265 -----YGEPMFKGEYKXNVEYPLYIQVRCFFELKKDKIPCIQIKGNARFGQNEYLS 317

Query: 414 XXXXXXXXLRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTQGKLKKNINM 473
 L +T +D I+ + + I+E E+ +F+ + I

Sbjct: 318 TSGDEYVDLY----VTNVDWELIKKH-YDIFEEEFIGG--FMFKGF-----IGF 359

Query: 474 TSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYGIPALRSHFN--LFRLLDDNNELNYINI 531
 Y + N S E+ + +K++LN LYG A + LD+N L

Sbjct: 360 FDEYIDRFMEIKNSPDSSAEQSLQAKMLNLSLYGKFATNPDTGKVPYLDENGVLKFRKG 419

Query: 532 GYKNTERNILFST---FVTSRSLYNLLVFPQYLTESEIDNFIYCDTDSLYMKSVMKPLL 588
 K ER+ +++ F+T+ + N+L Q L FIY DTDS++++ + +

Sbjct: 420 ELK--ERDPVYTPMGCFITAYARENILSNAQKLYP-----RFIYADTDSIHVEGLGEVDA 472

Query: 589 NPSLFDPIALGKWDIE 604
 + DP LG WD E

Sbjct: 473 IKDVIDPKKLGWDHE 488

>gi|118851|sp|P06950|DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP2)
 >gi|75812|pir|ERBP2Z DNA-directed DNA polymerase (EC
 2.7.7.7) - phage PZA >gi|216051 (M11813) gene 2 product
 [Bacteriophage PZA] >gi|224741|prf|1112171E ORF 2
 [Bacteriophage PZA]
 Length = 572

Score = 45.3 bits (105), Expect = 0.002
 Identities = 98/461 (21%), Positives = 166/461 (35%), Gaps = 110/461 (23%)

Query: 198 QLKTDFNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQLTYIHNDVILGMCHIHYSDIFP 257
 ++ DF T+ D D + Y ++TP++ YI ND+ I+ + I

Sbjct: 129 KIAKDFKLTVLKGDIDYHKERPVG-----EITPDEYAYIKNDIQIAEALL----IQF 178

Query: 258 NFDYKNLTFSLNIMESYLNEMTR-----FQLLNQYQDIKISYTHYHFDNMFYDIKSF 312
 +++T + ++ + + T+ F L+ D ++ Y

Sbjct: 179 KQGLDRMTAGSDDLKGFKDIIITTKFKKVPPTLSLGLDKEVRYA----- 222

Query: 313 YRGLNMYNTKYINKLIDEPFSDINSSYPYVMYHEKIPTWLYFYEHYSEPTLIPT--F 370
 YRGG N ++ K I E D+NS YP MY +P Y EP +

Sbjct: 223 YRGGFTWLNDRFKEKEIGEGMV-FDVNSLYPAQMYSRLLP-----YGEPIVFEKGKYV 273

Query: 371 LDDDNYSFLYKID----KDVFNDDLIIKIKSRVLRQXXXXXXXXXXXXXXXXXXLRMI 425
 D+D + I K+ + + IK +SR +

Sbjct: 274 WDEDYPLHIQHIRCEFELKEGYIPTTIQIK-RSRFYKGNELKSSGGEIADLW----- 324

Query: 426 QDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTQGKLKKNINMTSPYDYHITDDI 485
 ++ +D + + + +Y EY F T G K+ I+ + I

Sbjct: 325 --VSND-LELMKEHYDLYNVEYISGLK-----FKATTGLPKDFIDKWTHTKTSEGA 375

Query: 486 NEHPYSNEEVMLSKVVLNGLYG-----IPALRSHFN--LFRLLDDNNELNYINGY 533
 + L+K++LN LYG +P L+ + L FRL G

Sbjct: 376 KQ-----LAKMLNLSLYGKFASNPDTGKVPYKENGALGFRL-----GE 415

Query: 534 KNTERNIL--FSTFVTSRSLYNLLVFPQYLTESEIDNFIYCDTDSLYMKSVMKPLLNS 591
 + T+ + F+T+ + Y + Q D IYCDTDS+++ P +

Sbjct: 416 EETKDPVYTPMGVFITAWARYTTITAAQACF-----DRIIYCDTDSIHLTGTETIPDVIKD 470

Query: 592 LFDPIALGKWDIENEQIDKMFVLNHHKKYAY-----EVNGKI 627
 + DP LG W E+ + L K Y EV+GK+

Sbjct: 471 IVDPKKLGWHAHES-TFKRAKYLRQKTYIQDIYMKEVDGKL 510

>gi|2435429 (AF012250) unassigned reading frame (possible DNA
 polymerase) [Physarum polycephalum]
 Length = 544

Score = 44.9 bits (104), Expect = 0.002
 Identities = 118/545 (21%), Positives = 206/545 (37%), Gaps = 104/545 (19%)

Query: 179 TSIATLGKKLLDGGYLTESQLKTDFNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQLTYI 238
 T + L K L D + T Q F N M Y + CF L P++ I

Sbjct: 62 TQLFNLLKSLQDSSFYTFKQ-----FTYQNM-----YSLEISCF--LYPKKKILI 105

Query: 239 HNDVILGMCHIHYSDIFPNFD-----YNKL--TFSLNIMESY-LNEMTRFQLLNQYQD 290
 D+ +I Y+D+ ++ YN++ +++NI Y L+ ++ +

Sbjct: 106 -KDLNFFSENIYNDVVKDYKLLAILYNEIQTAYNININRKYLSTASLSLRFKKSFP 164

Query: 291 IKISYTHYHFDHMFYDIKSFYRGGLNMYNTKYINKLIDPCFSIDINSSYPYVMYHEK 350
K + D + +YI+ Y GG N I + + + + D+NS YPY+M EK
Sbjct: 165 EKYRLIPHLTRDED--NYIRKSYIGGRNE-----IFEHVAQRNYFYDVNSLYPYIMKKEK 217

Query: 351 IPTWLYFYEYSEPTLIPTFLDD-DNYFS----LYKIDKDVFNDDL---IKIKSRVLRQ 402
+P + Y + + F + +N+F L I+K N +L + IK+ V
Sbjct: 218 MPIGI---PEYRDKEYMKKFEKNIENFFGFIDVLITIEKTNNIPVLPYRMGIKNNV-EV 273

Query: 403 MXXXXXXXXXXXXXXXXXLRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
L + Q I+ IY + +++F+ Y +
Sbjct: 274 GIIYAGTGLRGIYFSEEIKLALKQGYKIIE-----IYSAYEYKEKEVVFEEYVEQ 323

Query: 463 TQGK-LKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG-----IPALRS 513
+ LK K D + D L K +LN LYG I +
Sbjct: 324 MYNRRLKAK-----DPALKD-----LYKKLLNTLYGRFGLVYEQIDIISP 363

Query: 514 HFNLFRLLDDNNELNYINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYC 573
L + DN + + + + N + + + + F Y T + + IY
Sbjct: 364 EKEL--ITDNTYISHDTTEFIDITANTCYNNIAITSAITSYARIFMYNTILNYNLHVYI 421

Query: 574 DTDLYMKSVVKPLNPSLFDPIALGKWDIENEQIDKMFVLNHHKYAY-EVNGKIKIASA 632
DTD L++K+ P+ + +L +GK+ +E+ + F+ N K Y Y +N I
Sbjct: 422 DTDGLFLKN---PIPDIALTTSKEMGKFRLESINAEAHFIAN-KFYIYAPINSPIIYKFK 477

Query: 633 GIPK-----NAFDTSVDFETFVR---EQFFDGAIIENNKSIYNEQGT-----ISYPSK 678
GIP N D + + +F +I NN Y+ Q + I Y +
Sbjct: 478 GIPLQKPIFNIHDIITQHKILNITLGHYFTFSIRLNNNQYTSFQASRKRLIPNYKTT 537

Query: 679 TEIVC 683

I+C

Sbjct: 538 PWIIC 542

>gi|1084487|pir||S41618 DNA polymerase - slime mold (Physarum
polycephalum) >gi|509721|dbj|BAA06121.1| (D29637) DNA
polymerase [Physarum polycephalum]
Length = 547

Score = 44.9 bits (104), Expect = 0.002

Identities = 118/545 (21%), Positives = 206/545 (37%), Gaps = 104/545 (19%)

Query: 179 TSIATLGKLLDGGYLTESQLKTDFTNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQLTYI 238
T + L K L D + T Q F N M Y + CF L P++ I
Sbjct: 65 TQLFNLLKSLQDSSFYTFKQ-----FTYQNM-----YSLEISCF--LYPKKKILI 108

Query: 239 HNDVILGMCCHIHYSDIFPNFD-----YNKL--TFSLNIMESY-LNNEMTRFQLLNQYQD 290
D+ +I Y+D+ ++ YN++ ++NI Y L+ ++ +
Sbjct: 109 -KDLYNFFSENIYNDVVKDYKLLAILYNEIQYAININIRKYITLSTASLSLRFKKSFP 167

Query: 291 IKISYTHYHFDHMFYDIKSFYRGGLNMYNTKYINKLIDPCFSIDINSSYPYVMYHEK 350
K + D + +YI+ Y GG N I + + + + D+NS YPY+M EK
Sbjct: 168 EKYRLIPHLTRDED--NYIRKSYIGGRNE-----IFEHVAQRNYFYDVNSLYPYIMKKEK 220

Query: 351 IPTWLYFYEYSEPTLIPTFLDD-DNYFS----LYKIDKDVFNDDL---IKIKSRVLRQ 402
+P + Y + + F + +N+F L I+K N +L + IK+ V
Sbjct: 221 MPIGI---PEYRDKEYMKKFEKNIENFFGFIDVLITIEKTNNIPVLPYRMGIKNNV-EV 276

Query: 403 MXXXXXXXXXXXXXXXXXLRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
L + Q I+ IY + +++F+ Y +
Sbjct: 277 GIIYAGTGLRGIYFSEEIKLALKQGYKIIE-----IYSAYEYKEKEVVFEEYVEQ 326

Query: 463 TQGK-LKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG-----IPALRS 513
+ LK K D + D L K +LN LYG I +
Sbjct: 327 MYNRRLKAK-----DPALKD-----LYKKLLNTLYGRFGLVYEQIDIISP 366

Query: 514 HFNLFRLLDDNNELNYINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYC 573
L + DN + + + + N + + + + F Y T + + IY
Sbjct: 367 EKEL--ITDNTYISHDTTEFIDITANTCYNNIAITSAITSYARIFMYNTILNYNLHVYI 424

Query: 574 DTDLYMKSVVKPLNPSLFDPIALGKWDIENEQIDKMFVLNHHKYAY-EVNGKIKIASA 632
DTD L++K+ P+ + +L +GK+ +E+ + F+ N K Y Y +N I
Sbjct: 425 DTDGLFLKN---PIPDIALTTSKEMGKFRLESINAEAHFIAN-KFYIYAPINSPIIYKFK 480

Query: 633 GIPK-----NAFDTSVDFETFVR---EQFFDGAIIENNKSIYNEQGT-----ISYPSK 678

288

GIP N D + + +F +I NN Y+ Q + I Y +
Sbjct: 481 GIPLQKPIFNHDIITQHKKILNITLGHYFTFSIRLNNQTYSFQASRKRKLIPNYKTT 540
Query: 679 TEIVC 683
I+C
Sbjct: 541 PWIIC 545

>gi|4877819|gb|AAD31446.1| (AF133505) DNA polymerase [Neurospora
crassa]
Length = 1035

Score = 44.1 bits (102), Expect = 0.004
Identities = 36/172 (20%), Positives = 82/172 (46%), Gaps = 14/172 (8%)

Query: 521 DDNNELNYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSL YM 580
+ N EL + ++G K+ I ++ + + + + + + S Y DTDS+++
Sbjct: 817 EKQNYELLSYLDGEKDDGFIINSTSIAAATASWSRILMYKHIINSA-----YTDTDSIFV 870
Query: 581 KSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHHKKYAYEVNGKIKIASAGIPKNAFD 640
+ KPL + + + K + + I + + + K Y + GK++I GI KN +
Sbjct: 871 E---KPLDSAFIGEGCGKFKAEYNGQLIKRAIFISGKLYLLDFGGKLEIKCKGITKNKDN 927
Query: 641 TSVDFETFVREQFFDG---AIIENNKSIYNEQGTISIYPSKTEIVCGNVYDE 689
T+ + + E ++G + + E GT+++ K ++ G YD+
Sbjct: 928 TTHNLDINDFEALYNGESRVLFQERWGRSLELGTVTVKYQKYNLISG--YDK 977

>gi|461962|sp|P33537|DPOM_NEUCR PROBABLE DNA POLYMERASE
>gi|283351|pir||S26985 probable DNA-directed DNA
polymerase (EC 2.7.7.7) - Neurospora crassa
mitochondrion plasmid maranhar (SGC3)
>gi|578156|emb|CAA39046| (X55361) putative DNA
polymerase [Neurospora crassa]
Length = 1021

Score = 44.1 bits (102), Expect = 0.004
Identities = 36/172 (20%), Positives = 82/172 (46%), Gaps = 14/172 (8%)

Query: 521 DDNNELNYNIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSL YM 580
+ N EL + ++G K+ I ++ + + + + + + S Y DTDS+++
Sbjct: 815 EKQNYELLSYLDGEKDDGFIINSTSIAAATASWSRILMYKHIINSA-----YTDTDSIFV 868
Query: 581 KSVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHHKKYAYEVNGKIKIASAGIPKNAFD 640
+ KPL + + + K + + I + + + K Y + GK++I GI KN +
Sbjct: 869 E---KPLDSAFIGEGCGKFKAEYNGQLIKRAIFISGKLYLLDFGGKLEIKCKGITKNKDN 925
Query: 641 TSVDFETFVREQFFDG---AIIENNKSIYNEQGTISIYPSKTEIVCGNVYDE 689
T+ + + E ++G + + E GT+++ K ++ G YD+
Sbjct: 926 TTHNLDINDFEALYNGESRVLFQERWGRSLELGTVTVKYQKYNLISG--YDK 975

>gi|2499511|sp|Q12471|6P22_YEAST 6-PHOSPHOFRUCTO-2-KINASE 2
(PHOSPHOFRUCTOKINASE 2 II) (6PF-2-K 2)
>gi|2131162|pir||S61066 6-phosphofructo-2-kinase (EC
2.7.1.105) - yeast (Saccharomyces cerevisiae)
>gi|2131163|pir||S71026 6-phosphofructo-2-kinase (EC
2.7.1.105) - yeast (Saccharomyces cerevisiae)
>gi|1085116|emb|CAA62371| (X90861)
6-phosphofructo-2-kinase [Saccharomyces cerevisiae]
>gi|1420028|emb|CAA99157| (Z74878) ORF YOL136c
[Saccharomyces cerevisiae] >gi|1628439|emb|CAA64733|
(X95465) 6-phosphofructo-2-kinase [Saccharomyces
cerevisiae]
Length = 397

Score = 40.6 bits (93), Expect = 0.041
Identities = 48/208 (23%), Positives = 92/208 (44%), Gaps = 29/208 (13%)

Query: 175 MKTNTSIATLGKKLLDGGYLTESQLKTDNFNYTFDKDNDMNDSEAYDYAVKCFAKLTPEQ 234
++ S AT+ K LL L+ + + FN K+ND ++ +A++T ++
Sbjct: 139 IRRQISCATISKPLL----LSNTSSEDLFN----PKNNDKKET-----YARITLQK 181
Query: 235 LTY-IHNDVIIIGMCHIHYSIDFPNFDYKLTFSNLNIMESYLNEMTRFQLLN---QYQD 290
L + I+ND +G+ S I + F + S+ +E++ F L+ Q
Sbjct: 182 LFHEINNDECDVGIFDATNSTI-----ERRRFIFEEVCSFNTDELSSPNLVPIILQVSC 235

289

Query: 291 IKISYTHYHFHDMNFY-DYIKSFYRGGLNMYNTKYINKLIDPCFSID-INSSYPYVMYH 348
 S+ Y+ H+ +F DY+ Y + + + + FS+D N + Y+ H
 Sbjct: 236 FNRFSIKYNIHNKSFNEDYLDKPYELAIKDFAKRLKHYYSQFTPFSLDEFNQIHRYISQH 295

Query: 349 EKIPTWLYFYEHYSEPTLIPTFLDDDDNY 376
 E+I T L+F+ + + P L+ +Y
 Sbjct: 296 EEIDTSLFFFNVINAGVVEPHSLNQSHY 323

>gi|2258375|gb|AAD11909.1| (AF007261) transcription initiation
 factor sigma [Reclinomonas americana]
 Length = 532

Score = 39.9 bits (91), Expect = 0.070
 Identities = 49/205 (23%), Positives = 84/205 (40%), Gaps = 14/205 (6%)

Query: 100 NHFLKDTMRYPDNITRENIYLSAEENEHTLMKEATILAKNQNVIL---EKRVKSSIN 156
 N+ + + F + ++IY+ + +KE L K NVI+ K +K N
 Sbjct: 177 NYLVKNSYLNLFKTVPHDSIYMNYSYIQTPNLILKEYLQLIKIINVILQINKNIKKGN 236

Query: 157 LDLTMLNGFKFNIIDNFM---KTNTSIATLGKKLLDGGYLTESQLKTDNFNYTIFDKDND 213
 L++++FL F + N++ K + + + K L Y+T L T Y K
 Sbjct: 237 LNISFLYKFYQELKWNYYIFINKISRNTQKINIKTLKNSYITFYNLITFIQYTTTKQRL 296

Query: 214 MNDSEAYDYAVKCFK--LTPEQLTYIHNDVILGMCHIHYSDFPNFDYN-KLTFSLNI 270
 D +K F K P+ +N +I G+ HI+ + N K+T I
 Sbjct: 297 KMDIFYKQIFIKTFLKQHKIPKINKIKNSLIKYGLTHIYDMILISILRENKIVTLKNRI 356

Query: 271 MESYLNNEMTFRQLLNQYQDIKISY 295
 + Y+ T + QY +KI Y
 Sbjct: 357 IFNMPYITT---ISKQY--VKIGY 376

>gi|15734|emb|CAA37450| (X53370) DNA polymerase (AA 1-575)
 [Bacteriophage phi-29]
 Length = 575

Score = 39.5 bits (90), Expect = 0.092
 Identities = 41/150 (27%), Positives = 64/150 (42%), Gaps = 36/150 (24%)

Query: 497 LSKVVLNGLYG-----IPALRSHFNL-FRLDDNNELYNIINGYKNTERNIL--F 542
 L+K++LN LYG +P L+ + L FRL G + T+ +
 Sbjct: 381 LAKLMLNSLYGKFASNPDTVKGVPYKENGALGFRL-----GEEETKDPVYTPM 429

Query: 543 STFVTSRSLYNLLVPFQYLTESEIDDNFIYCDTDSLYMKSVVKPLNPSLFDPIALGKWD 602
 F+T+ + Y + Q D IYCDTDS+++ P + + DP LG W
 Sbjct: 430 GVFITAWARYTTITAAQACY-----DRIIYCDTDSIHLTGTEIPDVIKDIVDPKLGYYA 484

Query: 603 IENEQIDKMFVLNHHKYAY----EVNGKI 627
 E+ ++ L K Y EV+GK+
 Sbjct: 485 HES-TFKRVKYLKQKTYIQDIYMKEVDGKL 513

Query= pt|110872 44AHJDORF002 Phage 44AHJD ORF |3789-5732|3 1
 (647 letters)

>gi|135273|sp|P27622|TAGC_BACSU TEICHOIC ACID BIOSYNTHESIS PROTEIN C
 >gi|478126|pir||D49757 techoic acid biosynthesis protein
 tagC - Bacillus subtilis (strain.168) >gi|143727
 (M57497) putative [Bacillus subtilis]
 >gi|2636103|emb|CAB15594.1| (Z99122) alternate gene
 name: dinC [Bacillus subtilis]
 Length = 442

Score = 112 bits (278), Expect = 7e-24
 Identities = 91/314 (28%), Positives = 147/314 (45%), Gaps = 58/314 (18%)

Query: 152 FELNELEPKFVMGFGGIRNAVNSQINIDKETNHMYSTQSDS----QKPEGFWINKLTPSG 207
 F+ + PK V QS N D++ + +Y+TQ S + + I +L+ G
 Sbjct: 7 P DFTNITPKLFTELRVADKTVLQSFNFDEKNHQIYTTQVASGLKQNTQSYRITRLSLEG 66

Query: 208 DLISSMRIVQGGHGTITIGLERQSNNGEMKIWLHHD----GVAKLLQVAYKDNVYVLDLEA 262
 + SM + GGHGT IG+E + NG + IW +D ++L+ YK LD E +
 Sbjct: 67 LQLDSMLLKHHGGHGTNIGIENR-NGTIYIWSLYDKPNETDKSELVCFYPYKAGATLD-ENS 124

290

Query: 263 KGLTDYTPQSLLNKHTFTPLIDEANDKLILRFGDGTIQVRSRADVKNHIDNVEKEMTIDN 322
 K L ++ H TP +D N +L +R + D KN+ N ++ +TI N
 Sbjct: 125 KELQRFSNMPP--DHRVTPALDMKNRQLAIR-----QYDTKNN--NNKQWVTIFN 170

Query: 323 SE-----NNDN-----RWMQGIADVDDLYWLSGNSSVNSHVQIGKYSLTGQKI 367
 + N +N ++QG +D LYW +G+++ S+ + +
 Sbjct: 171 LDDAIANKNNPLYTINIPDELHYLQGFLLDDGYLYWYTGDTNSKSYPNL-----ITV 222

Query: 368 YDYPFKLSYQDGINFPRD-----NFKEPEGICIYTNPKTKRKSLLAMTNGGGGKRFH 420
 +D K+ Q I +D NF+EPEGIC+YTNP+T KSL++ +T+G G R
 Sbjct: 223 FDSDNKIVLQKEITVGKDLSTRYENNFREPEGICMYTNPETGAKSLMVGITSGKEGNRIS 282

Query: 421 NLYGFFQLGEYEHF 434
 +Y + YE+F
 Sbjct: 283 RIYAYH---SYENF 293

>gi|142847 (M64050) DNase inhibitor [Bacillus subtilis]
 Length = 125

Score = 51.9 bits (122), Expect = 1e-05
 Identities = 35/116 (30%), Positives = 55/116 (47%), Gaps = 10/116 (8%)

Query: 152 FELNELEPKFVMFGGIRNAVNQISINIDKETNHMYSTQSDS----QKPEGFWINKLTPSG 207
 F+ + PK V QS N D++ + +Y+TQ S + + I +L+ G
 Sbjct: 7 FDFTNITPKLFTELRVADKTVLQSFNFDEKNHQIYTTQVASGLGKONTQSYRITRLSLEG 66

Query: 208 DLISSMRIVQGGHGTITGLERQSNEMKIWLHHD-----GVAKLLQVAYKDNVYVLD 258
 + SM + GGHGT IG+E + NG + IW +D ++L+ YK LD
 Sbjct: 67 LQLDSMLLKHHGGHTNIGMENR-NGTIYIWSLYDKPNETDKSELVCFPYKAGATLD 121

>gi|4038407 (AF103943) factor C protein precursor [Streptomyces
 griseus]
 Length = 324

Score = 39.1 bits (89), Expect = 0.10
 Identities = 61/269 (22%), Positives = 102/269 (37%), Gaps = 33/269 (12%)

Query: 172 VNQISINIDKETNHMYSTQSDSQKPEG---FWINKLTPSGDLISSMRIVQGGHGTITGLER 228
 V QS D ++ Q S P+ I +L SG+ + M ++ GHG +IG +
 Sbjct: 66 VQQSFTFDIVNRRLFVAQLKSGSPDDSGDLCTITQLDFSGNKLGHMYLLGFGHGVSIGAQ- 124

Query: 229 QSNEMKIWLHHDGVAKLLQVAYKDNVYVLDLEAKGLTDYTPQSLLNKHTFTP----- 281
 + +W D + + + + G T S L K H P
 Sbjct: 125 PVGADTYLWTEVD-----VNSNARGTRLARFKWNGATLSRTSSALAKHQVPVPGATEMTC 179

Query: 282 LIDEANDKLILRFGDGTIQVRSRADVKNHIDNVEKEMTIDNSENNDNRWMQGIADVDDDL 341
 ID N+++ +R+ + + +V + V + D QG A+ G +
 Sbjct: 180 AIDPVNNRMAIRYLTAAGRRYGIYNVADIAAGVYDKPLSDVPHPTGLGTGQYALYGSYV 239

Query: 342 YWLSGN-----SSVNSHVQIGKYSLTGQKIYDYPFKLSYQDGINFPRDNFKEPEGIC 394
 Y L+GN + NS+V + TG + + + G F+EPEG+
 Sbjct: 240 YQLTGNPYGPDNPNPGNSYVS--SVDVNTGALVQ----RAFTRAGSTL---TFREPEGMG 290

Query: 395 IYTNPKTKRKSLLAMTNGGGGKRFHNL 423
 IY + + L L +G G R NL+
 Sbjct: 291 IYRTAAGEVR-LFLGFASGVAGDRRSNLF 318

Query= pt|110873 44AHJDORF003 Phage 44AHJD ORF |6626-8389|2 1
 (587 letters)

>gi|138123|sp|P04331|VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9)
 >gi|75850|pir||WMBPT9 gene 9 protein - phage phi-29
 >gi|215327 (M14782) tail protein [Bacteriophage phi-29]
 >gi|225364|prf||1301270D gene 9 [Bacillus sp.]
 Length = 599

Score = 92.4 bits (226), Expect = 8e-18
 Identities = 126/618 (20%), Positives = 251/618 (40%), Gaps = 71/618 (11%)

Query: 5 TNFKFFYNTPTFT-DYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPY-NFIRDMEINVD 62
 TN + + PF+ DY+NT F S+ + ++F R + + SK + F ++ ++V
 Sbjct: 9 TNVRILADVFPFSNDYKNTRWFTSSSNQYNWF--NRKSRVYEMSKVTFMGFRENKPYVSVS 66

291

Query: 63 MQWHDAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLEQL 121
 + +Y+ F + D+ ++ +YAFV ++E+ N V ++F ID + T+ ++
 Sbjct: 67 LPIDKLYSASYIMFQNADYGNKWFYAFVTELEFKNSAVTYVHFIDVLQTMFDMKFQES 126

Query: 122 SNVNIERQHLSKRTYNYMLPMLRNDDVLKVSNNKYVYNQMQQYLENLVLFQSSADLSKK 181
 I R+H+ K + P + D+ L ++ + + + ++F S
 Sbjct: 127 F---IVREHV-KLWNDDGTPINTIDEGLSYGSEYDIVSVENHKPYDDMMFLVVISKSIM 182

Query: 182 FGT--KKEPNLDTSKGTIYDNITSPVNLVMEYGDFFINFMKMSAYPWITQNFQK---V 235
 GT ++E L+ ++ + + P+ Y+ + + D +I N V
 Sbjct: 183 HGTGGEESRLNDINASL-NGMPQPLCYIHPF-----YKDGKVPKTYIGDNNANLSPIV 236

Query: 236 QMLPKDFINTKQLEDVKTSEKITGLKTLKQGGKSKEWSLK-DLSL-----SFSNLQ 285
 ML F + D+ + +T LK K+ + LK D + N+
 Sbjct: 237 NMLTNIFSQKSAVNDI-VNMYVTDYIGLKLDYKNGDKELKLDKDMFEQAGIADDKHGNVD 295

Query: 286 EMMLSK-----KDEFKHMIRNEYMTIEFYDWNNGNTMLLDAGKISQK 326
 + + K KD+ ++ Y E D+ GN M L I+
 Sbjct: 296 TIFVKKIPDYEALEIDTGDKNWGGFTKDQESKLMMPYCVTEITDFKGNHMLKTEYINNS 355

Query: 327 TGVKLRTKSIIGYHNEVRVYPVDYNSAENDRPILAKNKEILIDTGSFLNTNITFNSFAQV 386
 +K++ + +G N+V DYN+ D + N+ S +N N
 Sbjct: 356 K-LKIQVRGSLGVSNNKVAYSVQDYNA---DSALSGGNRLTASLDSSLINNNPN----- 404

Query: 387 PILINNGILGQSQQANRQ--KNAESQLITNRIDNVLNG---SDPKSRFYDAVSASNLSP 441
 I I N L Q N+ +N +S ++ N I ++ G + + A+ +AS++
 Sbjct: 405 DIAILNDYLSAYLQGNKNSLENQKSSILFNGIMGIGGISAGASAAGGSALGMASV-- 462

Query: 442 TALFGKFNEEYNYFYKQQAEYKDLALQPPSVTSEMGNAFQIANSINGLTMKISVPSPE 501
 T + + QA+ D+A PP +T+ AF N G+ + +
 Sbjct: 463 TGMTSTAGNAVLQMAMQAKQADIANIPPQLTKMGNTAFDYGNGYRGVYVIKKQLKAEY 522

Query: 502 ITFLQKYMYLFGFEVNDYNSFIEPINSMTVCNYLKTGTYYTIRIDPMLMEQLKAILSEG 561
 L ++ +G+++N + + NY++ + DI+ +++++ I ++G
 Sbjct: 523 RRSLSFFHKYGYKINRVKK--PNLRTRKAFNYVQTKDCFISGDINNNDLQEIRTIFDNG 580

Query: 562 VRFWHDGSGNPMQLQNPL 579
 + WH D GN ++N L
 Sbjct: 581 ITLWHTDNIGNYSVENEL 598

>gi|138124|sp|P07534|VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9)
 >gi|75849|pir|WMBP9Z gene 9 protein - phage PZA
 >gi|216058 (M11813) tail protein (Bacteriophage PZA)
 Length = 599

Score = 81.9 bits (199), Expect = 1e-14
 Identities = 127/618 (20%), Positives = 248/618 (39%), Gaps = 71/618 (11%)

Query: 5 TNFKFFYNTPTFT-DYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQFPYNFIRDRME-INVD 62
 TN + + PF+ DY+NT F S+ + ++F + + SK + R+ I+V
 Sbjct: 9 TNVRILADVFPFSNDYKNTRWFTSSSNQYNWF--NSKTRVYEMSKVTFQGFRENKSYISVS 66

Query: 63 MQWHDAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLEQL 121
 ++ +Y+ F + D+ ++ +YAFV ++EY N ++F ID + T+ N+ Q
 Sbjct: 67 LRLDLLYNASYIMFQNADYGNKWFYAFVTELEYKNVGTYYVHFIDVLQTW-MFNIKPFQE 125

Query: 122 SNVNIERQHLSKRTYNYMLPMLRNDDVLKVSNNKYVYN--QMQQYLENLVLFQSSADLS 179
 S I R+H+ K + P + D+ L ++ + + + Y + + L S +
 Sbjct: 126 SF--IVREHV-KLWNDDGTPINTIDEGLNYGSEYDIVSVENHRPYDDMMFLVVISKSIM 182

Query: 180 KKGTKKEPNLDTSKGTIYDNITSPVNLVMEY-----GD-----FINFMDK 221
 + E L+ ++ + + P+ Y+ + GD +N +
 Sbjct: 183 HGTAGEAESRLNDINASL-NGMPQPLCYIHPFYKDGKVPKTFIGDNNANLSPIVNMLTN 241

Query: 222 MSAYPWITQNFQKQMLPKDFINTK-----DLEDVKTSEKITGLKTLKQGGKSKEWS 273
 + + N V M D+I K +L+ K + G+ K G +
 Sbjct: 242 IFSQKSAVNNI--VNMYVTDYIGLKLDYKNGDKELKLDKDMFEQAGIADDKHGNVDITIF 299

Query: 274 LKDL---SLSPSNLQEMMLSKKDEFKHMIRNEYMTIEFYDWNNGNTMLLDAGKISQKTGVK 330
 K +L + KD+ ++ Y E D+ GN M L I +K
 Sbjct: 300 KKIPDYETLEIDTGDKNWGGFTKDQESKLMMPYCVTEVTDFKGNHMLKTEYIDNNK-LK 358

Query: 331 LRTKSIIGYHNEVRVYPVDYNSAENDRPILAKNKEILIDTGSFLNTNITFNSFAQVPILI 390
 ++ + +G N+V DYN+ + L+ + L+T++ N+ + I+

292

Sbjct: 359 IQVRGSLGVSNKVAYSIQDYNAGGS----LSGGDRLTAS----LDTSLINNNPNDIAII- 409

Query: 391 NNGILGQSQQANRQ--KNAESQLITNRIDNVNLNGSDPKSRFYDAVSASNLSP----- 441
N L Q N+ +N +S ++ N I +L G A + A SP

Sbjct: 410 -NDYLSAYLQGNKNSLENQKSSILFNGIVGMLGGG-----VSAGASAVGRSPFGLASSV 462

Query: 442 TALFGKFNEEYNFYKQQQAQAEYKDLALQPPSVTESEMGNAPQIANSINGLTMKISVSPSKE 501
T + + QA+ D+A PP +T+ AF N G+ + +

Sbjct: 463 TGMTSTAGNAVLDQMALQAKQADIANIPQLTKMGNTAFDYGNGYRGVYVIKQLKAEY 522

Query: 502 ITFLQKYMYLFGFEVNDYNSFIEPINSMTVCNYLKTGTGTYTIRDIDPMLMEQLKAILES 561
L ++ +G+++N + + NY++ + DI+ +++++ I ++G

Sbjct: 523 RRSLSPPFHKYGYKINRVKK--PNLRTRKAYNYIQTKDCFISGDINNNDLQEIIRTIFDNG 580

Query: 562 VRFWHDGSGNPMLQNPL 579
+ WH D GN ++N L

Sbjct: 581 ITLWHTDDIGNYSVENEL 598

>gi|1429238|emb|CAA67657| (X99260) tail protein [Bacteriophage B103]
Length = 598

Score = 77.6 bits (188), Expect = 2e-13
Identities = 130/623 (20%), Positives = 240/623 (37%), Gaps = 86/623 (13%)

Query: 5 TNFKFFYNTPTT-DYQNTIHPNSNKERDDYFLNGRHFKSLDYSKQPYNFI---RDRMEIN 60
T+ + F N PP+ DY++T F + + YF + K + NF+ I

Sbjct: 9 TDVRFISNVFPFSNDYKSTRWFTNADAQYSYF---NAKPRVHVINECNFVGLKEGTPHIR 64

Query: 61 VDMQWHDAGQINMYTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLE 119
V+ + D YM F + + ++ +Y FV ++EYVN V +YF ID I T+ +

Sbjct: 65 VNKRIDDLYNACYMIFRNTQYSNKNWFYCFVTRLEYVNSGVNTNLYFEIDVIQTW-MFDFKP 123

Query: 120 QLSNVNIERQHLSKRTYNMPLMRNDDVLKVSNNKYVYNMQQYLENLVLFSQSSADLS 179
Q S + E Q + P+ D+ L + V Q ++F S

Sbjct: 124 QPSYIVREHQEMWDANNE---PLTNTIDEGLNYGTEYDVVAVEQYKPYGDLMPMVCISKS 180

Query: 180 KKGFTKKEPNLDTSGKITYDNITS---PVLNLYVMEYGDFFNFMKMSAYPHWITQNFQKVQ 236
K T E G I NI P++ YV + + D S P +T +VQ

Sbjct: 181 KMHATAGET---FKAGEIAANINGAPQPLSYVHPF-----YEDGSS--PKVTIGSNEVQ 230

Query: 237 ML-PKDFINTKQLEDVKTSEKITGLKT-----LKQGGKSKESLKDLSLSFSNL----- 284
+ P DF+ ++ + ++ T + +K SL+D + +

Sbjct: 231 VSKPTDPLKNMFTQEHAVNNIVSLYVTDYIGLNIHYDESATKMSLRDTMFEHAQIADDKH 290

Query: 285 -----QEMMLSKKDEPKHMIRNEYMTIEFY-----DWNQNTMLLDAGK 322
+E + +F NE + Y D+ GN + +

Sbjct: 291 PNVNTIYLKEVKEYEKTIDTGYKFASFANNEQSKLLMPYCVTTITDFKGNQIDIKNEY 350

Query: 323 ISQKTGVKLRTKSIIGYHNEVRVYPVDYNS---AENDRPILAKNKEILIDTGSFLNTNIT 379
++ + +K++ + +G N+V DYN+ D+ + A NT++

Sbjct: 351 VNG-SNLKIQVRGSLGVSNKVTVSVQDYNADTTLSGDQNLTA-----CNTSLI 398

Query: 380 FNSFAQVPILINNGILGQSQQANRQ--KNAESQLITNRIDNVNL---GSDPKSRFYDAVS 434
N+ V I+ N L Q N+ +N + ++ N + ++L G+ + AV

Sbjct: 399 NNNPNDVAII--NDYLSAYLQGNKNSLENQKDSILFNGVMSMLGNGIGAVGSAATGSAVG 456

Query: 435 VASNLSPALFGKFNEEYNFYKQQQAQAEYKDLALQPPSVTESEMGNAPQIANSINGLTMKI 494
VAS S T + + QA+ D+A PP + + A+ N G+ +

Sbjct: 457 VAS--SATGMVSSAGNAVLQIQGMQAKQADIANIPQLVKMGNTAYDYGNGYRGVYVIK 514

Query: 495 SVSPSKEITFLQKYMYLFGFEVNDYNSFIEPINSMTVCNYLKTGTGTYTIRDIDPMLMEQL 554
+ L + +G++ N + + + NY++ I +++ +++++

Sbjct: 515 KQIKEEYRNILSDFSRKYGYKTNLVK--MPNLRTRESYNYVQTKDCNIIGNLNEDLQKI 572

Query: 555 KAILESQVRFWHDGSGNPMLQN 577
+ I +SG+ WH D G+ L N

Sbjct: 573 RTIFDSGITLWHADPVGDTTLNN 595

>gi|215339 (M12456) p9 tail protein [Bacteriophage phi-29]
>gi|224163|prf||1011232C protein p9,tail [Bacteriophage
phi-29]
Length = 335

293

Score = 71.0 bits (171), Expect = 2e-11
 Identities = 64/293 (21%), Positives = 123/293 (41%), Gaps = 20/293 (6%)

Query: 292 KDEFKHMIRNEYMTIEFYDWNNGNTMLLDAGKISQKTGVKLRKTSIIIGYHNEVRVYPVDYN 351
 KD+ ++ Y E D+ GN M L I+ +K++ + +G N+V DYN
 Sbjct: 57 KDQESKLMMYPYCITEITDFKGNHMLKTEYINNSK-LKIQVRGSLGVSNNKVAISVQDYN 115

Query: 352 SAENDRPILAKNKEILIDTGSFLNTNITFNSFAQVPILINNGILGQSQANRQ--KNAES 409
 + D + N+ S +N N I I N L Q N+ +N +S
 Sbjct: 116 A---DSALSGGNRLTASLDSSLINNNPN-----DIAILNDYLSAYLQGNKNSLENQKS 165

Query: 410 QLITNRIDNVLNG--SDPKSRFYDAVSASNLSPALFGKFNEEYNFYKQQQAEYKDLA 466
 ++ N I ++ G + + A+ +AS++ T + + QA+ D+A
 Sbjct: 166 SILFNGIMGMIGGGISAGASAAGGSALGMASV--TGMTSTAGNAVLMQAMQAKQADIA 223

Query: 467 LQPPSVTESEMGNAFQIANSINGLTMKISVSPKEITFLQKYMLFGFEVNDYNSFIEPI 526
 PP +T+ AF N G+ + + L ++ +G+++N +
 Sbjct: 224 NIPPQLTKMGNTAFDYGNNGYRGVYVIKKQLKAEYRRSLSSPFHKYGYKINRVKK--PNL 281

Query: 527 NSMTVCNLYKCTGTYYTIRIDPMLMEQLKAIKESGVRFWHNDGSGNPMLQNPL 579
 + NY++ + DI+ +++++ I ++G+ WH D GN ++N L
 Sbjct: 282 RTRKAFNYVQTKDCFISGDINNNDLQEI RTIFDNGITLWHTDNIGNYSVENEL 334

>gi|1181968|emb|CAA87738.1| (Z47794) tail protein [Bacteriophage
 CP-1]
 Length = 230

Score = 53.9 bits (127), Expect = 3e-06
 Identities = 29/113 (25%), Positives = 54/113 (47%), Gaps = 3/113 (2%)

Query: 1 MRKLTNFKFFYNTPF-TDYQNTIHFNSNKERDDYFLNGRHFKSLDYKQKQPNFIRDRMEI 59
 M++ T + +PF DY N I+F + + +D+F + Y + + + I
 Sbjct: 1 MQESTKIWLYAKSPFKNDYANVINFETRESMEDFFTKKNPHIEIVYEYDKFYQTQRNGSI 60

Query: 60 NVDMQWHAQAQGINYMTFLSDFEDRRYYAFVNOQIEYVNDVVVKIYFVIDTIMTY 112
 V + + + YM F+++ R YYAFV + Y+N+ +I + +D TY
 Sbjct: 61 VVSGRVEKYENVTYMRFINN--GRYYAFVFDVLYINEDATRIIYEVDVWNTY 111

>gi|1181970|emb|CAA87740.1| (Z47794) tail protein [Bacteriophage
 CP-1]
 Length = 586

Score = 42.2 bits (97), Expect = 0.010
 Identities = 79/381 (20%), Positives = 139/381 (35%), Gaps = 92/381 (24%)

Query: 277 LSLSPSNLQEMMLSK--KDEFK---HMIRNEYMTIEFYDWNNGNTMLLDAG---KISQKT 327
 L +++ +QE + S KD+ + ++ +E+ IE YD GN+ + I +
 Sbjct: 187 LKIAVDQIQEGLRYSYMGKDDLEIEVQLLNSEFTEIELYDIYGNISYVYQPYLPRIDEAH 246

Query: 328 GVKLRKTSIIIGYHNEVRVYPVDYNSAEN----DRPIL----- 360
 K+ +G N+V + ++YN+A N D+ IL
 Sbjct: 247 KYKIVIVSGSLGDSNQVHINFLEYNNANNVSYADKNILDSLESGDWAHNPHEFKYGLNDV 306

Query: 361 -AKNKEILIDT-GSFLNTNITFNSFAQVPILINNGILGQSQANRQKNAESQLITNRIDN 418
 K+ IL D S++ ++ Q+ N +L QS + ++ A + +
 Sbjct: 307 TGKSVAILNDAEASYIQSHKNQMEHTQLTFKENRDMKQSVDLNKKQVATANSQASYNQA 366

Query: 419 VLNGSDPKSRFYDAVSASNLSPALFGKF-----NEEYNFYKQQQ-- 459
 S +++ + S N++ L G F N +YN QQ
 Sbjct: 367 FAVDSANINQWTEGASGILNVAGNLLTGNFGGALGLASGGMKVFNANRDYNDKVVQQGF 426

Query: 460 -----AEYKDLALQPPSVTESEMGNAFQIANSIN 488
 A DL QP SV + AFQ N +
 Sbjct: 427 TSENNALKSQSNALANMKSIALDQSI RAYNATMADLQNPISVQQIGNDLAFQSGNRLT 486

Query: 489 GLTMKISVSPKEITFLQKYMLFGFEVNDY-NSFIEPINSMTVCNLYKCTGTYY--TIRD 545
 + K+S+ + + +Y +G VN + N + + S NY+K T+R
 Sbjct: 487 DVYKVKSLAQKEIMGRANEYIKCYGVLVNWFTNDALSVMRSRKRPNYIKMINVNLGTLR- 545

Query: 546 IDPMLMEQLKAIKESGVRFWH 566
 + M ++AI +SGVR W+
 Sbjct: 546 ANQSHMNAIQAIQSGVRIWN 566

Query= pt|110875 44AHJDORF005 Phage 44AHJD ORF |12643-13890|-1 1
(415 letters)

>gi|3845203 (AE001399) GAF domain protein (cyclic nt signal
transduct.) [Plasmodium falciparum]
Length = 1245

Score = 52.3 bits (123), Expect = 6e-06
Identities = 59/246 (23%), Positives = 105/246 (41%), Gaps = 27/246 (10%)

Query: 174 ESIDRNHGNVDYIGFPPKMFLLGNAVNFSSPILSNLNIYNLLQKHKMTSRLYKNIFLEMR 233
+S D N+ N + + N+V FS+ N IY++L N +YK + E+
Sbjct: 854 DSSDNNNNNNNNNNNNNNNNNNNNSVIFST----NEKIYDML-----NRDNIYKKVKKEIF 904

Query: 234 RNDYVNEKRNTAFNSNDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYIKTDDKYI-- 291
D + + + +N + M + N N ++N+ N+ N NGD Y KY
Sbjct: 905 EGDSSIKTMEKPNLTNKNYMNNDNIDNNNNNNNNNNIDNNNNNGDNIYNDLKKYLYN 964

Query: 292 KVMYNTTFMTNIIIVVPYTKQYEFCTKIR-DIDNHVTYLRDDMFYKENMERYYYNPSNLH 350
++N ++ + + + K E K+ I + L +F+K NM + + L+
Sbjct: 965 TSIFNKDLYVKHFVDIIMNKSLEEIIMNVYISERINSL---LFHKGNM---LNDVTKLY 1018

Query: 351 FDNAYSKNYVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFERKEKIYEDN----YIENTK 406
NAY + N K I F + E K +M F+ +KIY+ N + N K
Sbjct: 1019 MSNAYGEKCFFFN-----FPQIKEIIFVNEYEKMDMKYFKMLKKIYKYNLKNKIFSNNYK 1073

Query: 407 KYLMKQ 412
+++K+
Sbjct: 1074 FFIIKK 1079

>gi|3758843|emb|CAB11128.1| (Z98551) predicted using hexExon;
MAL3P6.23 (PFC0820w), Hypothetical protein, len: 4982 aa
[Plasmodium falciparum]
Length = 4981

Score = 49.2 bits (115), Expect = 5e-05
Identities = 67/287 (23%), Positives = 110/287 (37%), Gaps = 60/287 (20%)

Query: 127 ITDLNSATDLKYHSNFKLKHYPPIIYDEFLALEDYLDIEWDKLKT----IYESIDRNHGN 182
I D+N + D+ + +++ I YD +++DK++ IY +ID++ N
Sbjct: 3619 IMDINKSKDISKNMEIVQN---IEYD-----NKYDKIRNDMDAIYMAIDKMDN 3664

Query: 183 VDYIGFPPKMFLLGNAVNFSSPILSNLNIYNL---LQKHKMTSRLYKNIFLEMRNDYV 238
+ I + F L N S +N YNL ++ K N R Y N F +D
Sbjct: 3665 IGIINCRRYFNLYKNYNNLSNECNRE-YNLNELYMEDIKRNMKR-YDNFNINHYDDNN 3722

Query: 239 NEKRNTAFNSNDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYIKTDDKYIKVMYNT 298
N N N+N++ N N ++N N+ N NG F+ D
Sbjct: 3723 NNGGCGFFHVD----- 3771

Query: 299 TFMNTNIIIVVPYTKQYEFCTKIRDIDNHVTYLRDDMFYKENMERYYYNPSNLHFDNAYSKN 358
K FCTK ++F +N+E N N N Y+ N
Sbjct: 3772 -----KDLFFCTK-----KNIFPCKNIETVCKNEYNKKIYNNYTCN 3807

Query: 359 YVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFERKEK-IYEDNYIEN 404
V+N + ++IK + + N E+ + EK +Y + EN
Sbjct: 3808 ISVNTNLNCLNIIKELIKLNNKKILNYYEYHKVEKLLYYRHSPEN 3854

Score = 35.6 bits (80), Expect = 0.70
Identities = 62/290 (21%), Positives = 121/290 (41%), Gaps = 65/290 (22%)

Query: 2 VKQNRDLMDVRDYQNAV--HVRKKIPDKYNQIELVDELMDDIDYIISINRSDGKSFNY 59
+K+N ++ +N +N +V++ DK N I D++I+ SN + +SF
Sbjct: 4445 IKRNNINKSNIKRNNINKSNVKSNTDKSNVIS-----DFHIT-SNNNITRSFT- 4492

Query: 60 VSPFIYLAIKLDIKFTLLSRHYTLRDAYRDFIEEIIDENPLFKSKRVTFRSARDYLAIY 119
A D F LS TL +Y +F + + I
Sbjct: 4493 -----ATLTDISIFNTLSE--TLNYSYDNFFSNMDN-----IKI 4523

Query: 120 QDKEIGVITDLNSATDLKYHSNFKLKHYPPIIYDEFL----ALEDDYLDIEWDKLKTIE 174
+ EI ITD++ +YH N+LK + +E++ + +D + DE ++T+ E
Sbjct: 4524 KKEINNITDQVYGNKKEYHENYLVKQNKVNEEYIEETFKSDKDCSIKDEACTIRTLS 4583

295

Query: 175 S--IDRNHGNVDYIGFPMFLGNVNFSSPILSNLNIYNLLQKHKMN--TSRLYKNIFL 230
 S I N N+D + + + S P N++ N++K+ +N R+ KN
 Sbjct: 4584 SCNISENINID-----MDEHISFPNGRNVHDNNYMKKNHVNYDKMRVKGKIP 4634

Query: 231 EMRRNDYVNEKRNTAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGD 280
 D + +++ + +D M++ ++ E ++ + L + NG+
 Sbjct: 4635 SFTHFDKILDEKKKK----SDKDMSSSKWLEREEHIKEIKLEKNEYMNGN 4680

Score = 34.0 bits (76), Expect = 2.0
 Identities = 47/211 (22%), Positives = 84/211 (39%), Gaps = 32/211 (15%)

Query: 210 IYNLLQKHKMTSRLYKNIFLEMRRNDYVNEKRNTAFNSNDDAMTTGEFEFNEYNLADD 269
 I++LLQK LY+N+ + R + N+ T E ++ + ++
 Sbjct: 918 IFSLLQKDSPLLVLYENVHI-----REGEKYGRNE--ATDNEVDYKKGDIKH 964

Query: 270 NLRNHINQNGDFFYIKTD---DKYIKVMYNTTFTMTNIIIVPYTKQYEFCTKIRDIDNHV 326
 N+ N + D + D+ K MY + V E K D+ N+
 Sbjct: 965 NVTNEHGNHSDSYPGNSLNLDRKPKNMYE-DIYKEKGFVKSDCSNIEI--KKNDMINND 1021

Query: 327 TYLRDDMPYKENMERYYPNSNLHFDNAYSKNYVVDNDRYLYLDMNKII---KPHIKNE 382
 Y +++ FY+++ Y+ + YV++ +YL +N ++ F +KN+
 Sbjct: 1022 VYKQNE-PYEDSRINMIYDEDEIKTWFLIPKHYVIN--IIYFLNILLTDESNEFKLKNK 1077

Query: 383 MKQNMSEFERKEKIYEDN-----YIENTKKY 408
 E K IYEDN ++N KKY
 Sbjct: 1078 KYGVFNZETKGTIYEDNNGLBILKNGKKY 1108

Score = 33.6 bits (75), Expect = 2.7
 Identities = 42/198 (21%), Positives = 77/198 (38%), Gaps = 42/198 (21%)

Query: 222 SRLYKNIFLEMR---RNDYVNEKRNTAFNSNDDAMTTGEFEFNEYNLA 267
 S LY I++ + +N + K+NT + N+++D TT E + +
 Sbjct: 411 SVLYSIYMNKKYKKNFIITNKKNTNVVFENDVIQLSVENTSSEDFTTNTRESSLNSGM 470

Query: 268 DDNLRNHINQNGDFFYIKTDKDYIKVMYNTTFTMTNIIIVPYTKQYEFCTKIRDIDNHVT 327
 +++R +N D +DDK ++Y N YTK E
 Sbjct: 471 MNDMRYGVNNYADEKVYHSDDKSDHLIYKHVHDEKKNKYDEMYTKTKE----- 517

Query: 328 YLRDDMPYKENMERYYPNSNLHFDNAYSKNYVVDNDRYLYLDMNKIIKPHIKNEMKKNM 387
 +++ YK N+ + N K LD+ K I H+KN+ + N
 Sbjct: 518 --NENIYKSNIVDKKTCDISSEMVNGKDK-----LDVEKYIGSHVKNDENNK 563

Query: 388 SEFERK-EKIYEDNYIEN 404
 + ++K + + + YI+N
 Sbjct: 564 EKLKKKIDNVNKKYIDN 581

>gi|3845297 (AE001421) hypothetical protein [Plasmodium falciparum]
 Length = 2380

Score = 48.0 bits (112), Expect = 1e-04
 Identities = 87/390 (22%), Positives = 160/390 (40%), Gaps = 65/390 (16%)

Query: 20 VRKKIPDKYNQIELVDELNDDIDYISISNRSKGKSFNYVSFF-----IYLAIKLDIKF 74
 +++K +K ++ + +N D + ++ R K+ NY++ +YL I DI
 Sbjct: 1049 LQRKNMKNCKSKNRNRNRYINKDSNIHLMNLIRIKFQNLNVMNMSFEIELYLKINNDIFL 1108

Query: 75 TLLSRHYTLRDAYR-----DFIEEIIIDEN-PLFKSKRVTFRSARDYLAIYQDKIGVI 127
 +Y +++ Y + + + EN + +++ ++ + Y +K+
 Sbjct: 1109 QFNKGNYNVQNFYNSITLINIMSKYSENFYAYNLEKIVYKFLNNKNFEYIEKQYSSK 1168

Query: 128 TDLSATDLKYHNSFLKHYPIIIYDEFLA---LEDDYLIDEWDKLTIIYESIDRNHGNV 183
 D+N D+ ++ +K+ II EFL L+ D I + KLKT ++
 Sbjct: 1169 EDMNEL-DILVNTYDMKYDKII---EFLQNGYGLKIDRYIYFYPKLKT-----DI 1214

Query: 184 DYIGFPMFLGNVNFSSPILSNLNIYNLLQKHKMTSRLY-----KNIF--LEMRRN 235
 F ++FL N+ L NI +++ K + Y K IF + M+ +
 Sbjct: 1215 ILFFFKEIFLNDNLIKIDRKFLKK-NITIMIEVLKEIFKEYVKRCITKVIFPVMHKEH 1273

Query: 236 DYVNEKR-----NTRAFNSNDDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYIKTD 287
 D+V K N+ FN+ D + N YN D+ N+ N N +Y K

296

Sbjct: 1274 DHVMNKYNNQYVNNNSNMFNTRGDHNNNNQTNNDNHYNNHYDDTHNNNNNNNSKYK-KNK 1332

Query: 288 DKYIKVMYNVTFMTNIIV---VPYTKQYEFCTKIRDIDNHVTVLRDDMFYKEN---ME 340
+K K+MY +++ + V K + K I + Y+ ++ N +

Sbjct: 1333 NKN-KIMYKERKSSSLFISNNVQDVKPIKHYLYSSIIYKNFIYIIEIKNFNNKITKIN 1391

Query: 341 RY-YNPSNLHFDNAYSKNYVVDNDRYLYL 369
RY YYN NL+ D+ ND YL+L

Sbjct: 1392 RYNYNYMNLNIDDL-----NDAYLFL 1413

Score = 32.5 bits (72), Expect = 6.0
Identities = 46/183 (25%), Positives = 73/183 (39%), Gaps = 26/183 (14%)

Query: 225 YKNIFLEMRNDYVNEKRNTRAFNSNDDAMTGEFEFNEYNLADDNLRNHINQNGDFFYI 284
+KNI ++ ++N + NSN + + N N+ +N N IN + I

Sbjct: 27 HKNINKNIKKNKFINIDNSNCCNSNSNSNNNNNNNNNIVRNN--NNFINADKKKNVI 85

Query: 285 KTDDKYIKVMYNVTFMTNIIVVPYTKQYEFCTKIRDIDNHVTVLRDDMFYKENMERYYY 344
+D IK V NI Y ++ + D+ N+ + + KE ER

Sbjct: 86 LNEDDDIKNKELVDES FVNIPF--YENYFKNLFLNDVSNKVI--NIIEQKEGDER--- 138

Query: 345 NPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFERKEKIYEDNYIEN 404
N N N +KN V DN +NK IKN +N++E Y N++ +

Sbjct: 139 NADN----NLKNKNIVRDN-----INK-----IKN--TRNVNEILIYNKYIINFLND 180

Query: 405 TTK 407
T K

Sbjct: 181 TTK 183

>gi|4493936|emb|CAB38972.1| (AL034556) predicted using hexExon;
MAL3P5.6 (PFC0600w), Hypothetical protein, len: 250 aa
[Plasmodium falciparum]
Length = 249

Score = 47.3 bits (110), Expect = 2e-04
Identities = 53/215 (24%), Positives = 87/215 (39%), Gaps = 30/215 (13%)

Query: 209 NIYNLLQKHKNMNTSRLYKNIFLEMRNDYVNEKRNTRAFNSNDDAMTGEFEF--NEYNL 266
NIYN L++ YKN N ++ +N N+N EFE N YN

Sbjct: 13 NIYNKLEEK-----YKNFLKLKNMNSHMGASQNMNV--NNNYTMNELEEFKINNNYNN 64

Query: 267 ADDNLRNHINQNGDFFYIKTD-----DKYIKVMYNVTFMTNIIVVPYTKQYEFCTKIRD 321
++N+ N+IN D+ IK +K ++ YN + I T ++

Sbjct: 65 NNNNNNNNNNNYDYVMNIKVSQSVQHNRQLQDFYNNKNSFQHYIKLKTCTCFDADDIRNL 124

Query: 322 IDNHVTVLRDDMFYK-----ENMERYYYNPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIK 376
++ + Y RD+ K EN + N + N+ S NY DN+ LY +N++ K

Sbjct: 125 LEKRLAYERDNTLIKNIQEEENKKGIGINGNFGSESNSSSSNY--DNNYLYRKINRLNK 182

Query: 377 FHIKNEMKKNMSEFERKEKIYEDNYIENTKKYLMK 411
+ ++ KI KKY++K

Sbjct: 183 TNTNKSKNRSRKRKRINSKI-----DKKYIIK 209

>gi|3845165 (AE001390) hypothetical protein [Plasmodium falciparum]
Length = 1247

Score = 45.7 bits (106), Expect = 6e-04
Identities = 52/239 (21%), Positives = 94/239 (38%), Gaps = 38/239 (15%)

Query: 206 SNLNIYNLLQKHKNMNTSRLYKNIFLEMRNDYVNEKRNTRAFNSNDDAMTGEFEFNEYN 265
+N N +N ++K K R I +N + +N ++N+D E N N

Sbjct: 474 NNTNKNWEIKRKKKKFKREKNKIINNSFQNEAEDDKNNNNNDNNNDNHNNDNNNNEN 533

Query: 266 LADDNLRNHINQNGDFFYI-KTDDKYIK---VMYNVTFMTNIIVVPYTKQYEFCTKIR 320
D+N N+ + ND I D+ Y +YN T ++ YTK + + +

Sbjct: 534 NNDNNNNNNNDINNDINNHNNDNNYNNNDNINLYNEMTKKKMLDNSYTKYFFYIFTL- 592

Query: 321 DIDNHVTVLRDDMFYKENME-----RYYN-----PSNLHFDNAYS 356
+ + ++ + FY++N + ++YNN + N

Sbjct: 593 ---DMLPSIKFETFEKNTDHKNFNENYKFYYNTDDDTDIINAIKKNVKNKKKNGNIVI 649

Query: 357 KNYVVDNDRYLYLDMNKIIKFHIKNEMKKNMSEFER----KEKIYEDNYIENTKKYLMK 411
KNY+ N+ Y YL+ N+ + I + K +E K+ I+ ++Y E K K

297

Sbjct: 650 KNYINHNE-YSLEYENENKNEYEINKKEKLLTENYEYDMYIKDNIHYNDYSEGDKQTKK 707

Score = 41.0 bits (94), Expect = 0.016
Identities = 58/245 (23%), Positives = 96/245 (38%), Gaps = 43/245 (17%)

Query: 207 NLNIYNLLQKHKMTSRLYKIFLEMRRNDYVNEKRNTAFNSNDAMTTGEFEFNEYNL 266
N+N+YN + K K Y F + D + + + N D E YN
Sbjct: 564 NINLYNEMTKKCMLDNSYTKYFFYIFTLDMLPSIKFETFEKNTDHNKFNENYKFFYNT 623

Query: 267 ADD-----NLRNHINQNGDFF---YIKTDDKYIKVMYNVT-TFMTNIIIVVPYTKQ 312
DD N++N +NG+ YI ++ Y + YN + N T+
Sbjct: 624 DDDTDIINAIAKKKNVKNK-KKNGNIVIKNYINHNE-YSLEYENENKNEYEINKKEKLLTEN 681

Query: 313 YEFCTKIRDIDNHVTYLRDDMFYKENMERYYYNPSNLHFDNAYSK-----NYV--VD 362
YE+ I+D ++ Y D + + YN +N +N Y K +Y+ VD
Sbjct: 682 YEYDMYIKDNIHYNDYSEGDKQTKKASSFLYNNNN---NNKYKEDNKTQIISYMDHVD 738

Query: 363 NDR-----YLYLDMNKIIKFHIK-NEM---KQNMSEFERKEKIYEDNYIENTKKY 408
N+ Y + +++ F +K N+M K+ F +E I + +EN K+
Sbjct: 739 NENGKGLKKRNLFYNNSDQLYNFVDKNDMIKYEKQSKNFVEEEFINGNRKMENEDKH 798

Query: 409 LMKQY 413
L K Y
Sbjct: 799 LKKHY 803

Query= pt|110877 44AHJDORF007 Phage 44AHJD ORF |2044-3027|1 1
(327 letters)

>gi|1181960|emb|CAA87731.1| (Z47794) connector protein
[Bacteriophage CP-1]
Length = 337

Score = 45.7 bits (106), Expect = 5e-04
Identities = 44/184 (23%), Positives = 84/184 (44%), Gaps = 13/184 (7%)

Query: 127 QIHKLYDNCMSGNFVVMQNKPIQYNSDIEIIEHYTDELAVALSRFSLIMQAKFSK--IF 184
++HK + + +V+ N Y I +E + ++LA++ L+ L A+ + IF
Sbjct: 125 ELHKDNPKIKRCPVIPPNNF-YEPYIGYLELFCCKLADIELT-IQLNRNAQITPYFIF 182

Query: 185 KSEINDESINQLVSEIYNGAPFVKMSPMFNAD-----DDIIDLTSNSVIPALTEMKR 236
N S+ + ++I N P V ++ + D D I + L ++
Sbjct: 183 ADNTNVLMSKNIFNKIANFEPVVYLKQKQDQDQSFQKLSQDIQVFRTPDAPFLLDKLDH 242

Query: 237 EYQNKISELSNYLGINS LAVDKESGVSDDEAKSNRGFTTSNSNIYLGKREP-ITFLSKRY 295
E +++L ++GIN+ DK+ + EA SN G ++N + K R + ++K Y
Sbjct: 243 EKLVRMNQLLTFIGINNPSDKKERLVVSEAISNNGVISANIEVGWKSRRKFVELINKCY 302

Query: 296 GLDI 299
GL+I
Sbjct: 303 GLEI 306

>gi|1429239|emb|CAA67658| (X99260) upper collar protein
[Bacteriophage B103]
Length = 308

Score = 44.9 bits (104), Expect = 8e-04
Identities = 40/159 (25%), Positives = 73/159 (45%), Gaps = 11/159 (6%)

Query: 150 YNSDIEI-----IEHYTDELAEVA-LSRFSLIMQAKFSKIFKSEINDESINQLVSEIYNG 203
YN+D++ +E + +LAE+ + + Q I ++ N S+ + ++
Sbjct: 121 YNNDLKCSTLPALMFQAQLAELKEIIAVNQNAQKTPVLIANDNNQLSLKNIYNQYEGN 180

Query: 204 APFVKMSPMFNADD-DIIDLTSNSVIPALTEMKREYQNKISELSNYLGINS LAVDKESGV 262
AP + + + D+ + + V+ L K N E+ YLGI + ++K+ +
Sbjct: 181 APVIFVHESLDLDNLKVKFTDAPYVVDKLNQKNAVWN---EVMTYLGIKVANLEKKERM 237

Query: 263 SDEEAKSNRGFTTSNSNIYLGKGR-EPITFLSKRYGLDIK 300
E SN S+ NIYK R E +S+ YGL++K
Sbjct: 238 VTSEVDSNDEQIESGNIYKARQEACNKISELYGLNLK 276

>gi|137915|sp|P07535|VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR
PROTEIN) (LATE PROTEIN GP10) >gi|75851|pir||WMBP10 gene

298

10 protein - phage PZA >gi|216059 (M11813) upper collar
protein [Bacteriophage PZA]
Length = 309

Score = 43.8 bits (101), Expect = 0.002
Identities = 38/160 (23%), Positives = 75/160 (46%), Gaps = 13/160 (8%)

Query: 150 YNSDIEI-----IEHYTDELAELVALSRFSLIMQAKFSKIF--KSEINDESINQLVSEIYN 202
YN+D+ +E+ ELAE+ S+ A+ + + ++ N S+ Q+ ++
Sbjct: 122 YNNDMSPFTTPTTLELFAELAEELK-EIISVNQNAQKTPVLIRANDNNQLSLKQVYNQYEG 180

Query: 203 GAPFVKMSPMFNADD-DIIDLTNSVIPALTEMKREYQNKISELSNYLGINS LAVDKESG 261
AP + ++D ++ + V+ L K N E+ +LGI + ++K+
Sbjct: 181 NAPVIFAHEALDSDSIEVFKTDAPYVVDKLNQKNAVWN---EMMTFLGIKNANLEKKER 237

Query: 262 VSDEEAKSNRGFTTSNSNIYLKGR-EPITFLSKRYGLDIK 300
+ +E SN S+ ++LK R E +++ YGLD+K
Sbjct: 238 MVTDEVSSNDEQIESSGTVFLKSREEACEKINELYGLDVK 277

>gi|137914|sp|P04332|VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR
PROTEIN) (LATE PROTEIN GP10) >gi|75852|pir||WMBPC9 gene
10 protein - phage phi-29 >gi|215328 (M14782) upper
collar protein [Bacteriophage phi-29] >gi|215340
(M12456) p10 connector protein [Bacteriophage phi-29]
>gi|224161|prf||1011232A protein p10,connector
[Bacteriophage phi-29] >gi|225365|prf||1301270E gene 10
[Bacteriophage phi-29]
Length = 309

Score = 41.4 bits (95), Expect = 0.009
Identities = 37/160 (23%), Positives = 75/160 (46%), Gaps = 13/160 (8%)

Query: 150 YNSDIEI-----IEHYTDELAELVALSRFSLIMQAKFSKIF--KSEINDESINQLVSEIYN 202
YN+D+ +E+ ELAE+ S+ A+ + + ++ N S+ Q+ ++
Sbjct: 122 YNNDMAFPTTPTTLELFAELAEELK-EIISVNQNAQKTPVLIRANDNNQLSLKQVYNQYEG 180

Query: 203 GAPFVKMSPMFNADD-DIIDLTNSVIPALTEMKREYQNKISELSNYLGINS LAVDKESG 261
AP + ++D ++ + V+ L K N E+ +LGI + ++K+
Sbjct: 181 NAPVIFAHEALDSDSIEVFKTDAPYVVDKLNQKNAVWN---EMMTFLGIKNANLEKKER 237

Query: 262 VSDEEAKSNRGFTTSNSNIYLKGR-EPITFLSKRYGLDIK 300
+ +E SN S+ ++LK R E +++ YGL++K
Sbjct: 238 MVTDEVSSNDEQIESSGTVFLKSREEACEKINELYGLNVK 277

Query= pt|110878 44AHJDORF008 Phage 44AHJD ORF |3020-3775|2 1
(251 letters)

>gi|4982468|gb|AAD30963.2| (AF118151) SNF1/AMP-activated kinase
[Dictyostelium discoideum]
Length = 718

Score = 52.3 bits (123), Expect = 3e-06
Identities = 28/118 (23%), Positives = 56/118 (46%), Gaps = 5/118 (4%)

Query: 121 YLQSQGFTEHNEDTTSNTDETSQNATSLDNSTGMTANRNAYV---SLPQSEVNIDVDN 176
+ + GF N ++ SN + +N N + N+ T N N + ++ + +N + +N
Sbjct: 382 FTTTGFNPTNSNSISNNNNNNNNNNNTTNNNNNTTNNNNNSIINNNNNNNNNNNNNN 441

Query: 177 TTLRFADNNTIDNGKTVNKSSNESNQAKRNQKGNAGTQFTKQYLID-NIDKAYD 233
+NN I+N N ++N +N N N N N+ + T+ + I N++ +Y+
Sbjct: 442 NNNNNNNNNIINNNNNNNNNNNNNNNNNNNNNNSSISGGTEVFISPNLNNSYN 499

Score = 37.5 bits (85), Expect = 0.094
Identities = 17/111 (15%), Positives = 45/111 (40%)

Query: 130 HNEDTTSNTDETSQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189
+N + +N + +N N + +N++ ++ + P + + + + N+ ++
Sbjct: 456 NNNNNNNNNNNNNNNNNNNNNNNSSISGGTEVFISPNLNNSYNSSSGNSNGSNSNNNS 515

Query: 190 GKTIVNKSSNESNQAKRNQKGNAGTQFTKQYLIDNIDKAYDLRKKILN 240
N +N +N N N N N ID+++ + + + N
Sbjct: 516 NNNNTNNDNNNNNNNNNNNNNNNNNNNNNNNCIDSVNNSLNNENDVNN 566

Score = 31.7 bits (70), Expect = 5.4
Identities = 25/115 (21%), Positives = 48/115 (41%), Gaps = 10/115 (8%)

Score = 31.7 bits (70), Expect = 5.4
Identities = 15/104 (14%), Positives = 43/104 (40%)

Score = 30.9 bits (68), Expect = 9.2
Identities = 16/84 (19%), Positives = 34/84 (40%)

```
>gi|1730077|sp|P18160|KYK1_DICDI NON-RECEPTOR TYROSINE KINASE SPORE  
LYSIS A (TYROSINE-PROTEIN KINASE 1) >gi|974334 (U32174)  
non-receptor tyrosine kinase [Dictyostelium discoideum]  
Length = 1584
```

Score = 46.5 bits (108), Expect = 2e-04
Identities = 29/106 (27%), Positives = 48/106 (44%), Gaps = 4/106 (3%)

Score = 34.0 bits (76), Expect = 1.1
Identities = 20/117 (17%), Positives = 46/117 (39%)

Query: 87 NRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETSNQNA 146
N G I T + + + + + + + N + + N + + N N

300

Sbjct: 415 NNNNNNIIGNGKITTTTTSTSPSSINNNEDISSNNNNNNNNNNNNNNNNNNNN 474

Query: 147 TSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVNKSSNESQN 203
+ + + + + T N N + + N + + N N + + N N + + N + N N

Sbjct: 475 NNNNSNSSNTNNNNINNTTNNNSNSNNNNNNNSNSNSNNNNNNNNNNNNNNNN 531

Score = 33.2 bits (74), Expect = 1.8
Identities = 18/88 (20%), Positives = 35/88 (39%)

Query: 130 HNEDTTSNTDETSNQATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189
+ N + + + N + + N N T T + S + + E + N + N N + N

Sbjct: 405 NNNNSNNNNNNNNNNNNIIGNGKITTTTTSTSPSSINNNEDISSNNNNNNNNNNNNNN 464

Query: 190 GKTIVNKSSNESQNAKRQKQKGNKGT 217
N + + N + N N + + N T

Sbjct: 465 NNNNNNNNNNNNSNSSNTNNNNINNT 492

Score = 32.5 bits (72), Expect = 3.1
Identities = 18/94 (19%), Positives = 37/94 (39%)

Query: 120 KYLQSQGFTEHNEDTTSNTDETSNQATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTL 179
K + S N + + N + + N N + + + + T S N D + +

Sbjct: 392 KNVNSTSLVPGNNNNNNNNNNNNNNNNNNNNIIGNGKITTTTTSTSPSSINNNEDISSNN 451

Query: 180 RFADNNTIDNGKTVNKSSNESQNAKRQKQKGN 213
+ N N + N + + N + N N + + N

Sbjct: 452 NNNNNNNNNNNNNNNNNNNNNNNNNNSNSSNTN 485

Score = 32.5 bits (72), Expect = 3.1
Identities = 24/110 (21%), Positives = 44/110 (39%), Gaps = 10/110 (9%)

Query: 138 TDETSNQATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGK----- 191
T T + + + S + + N + + + N N + + N + N + N N

Sbjct: 429 TTTTTSTSPSSINNNEDISSNNNNNNNNNNNNNNNNNNNNNNNNNNNNNSNSSNTNNNN 488

Query: 192 ----TVNKSSNESQNAKRQKQKGNKGTQFTKQYLIDNIDKAYDLRKK 237
T N + SN + N N N N N + + N + L KK

Sbjct: 489 INNTTNNNSNSNNNNNNNSNSNSNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 538

>gi|3758855|emb|CAB11140.1| (Z98551) predicted using hexExon;
MAL3P6.11 (PFC0760c), Hypothetical protein, len: 3395 aa
[Plasmodium falciparum]
Length = 3394

Score = 46.5 bits (108), Expect = 2e-04
Identities = 52/202 (25%), Positives = 96/202 (46%), Gaps = 32/202 (15%)

Query: 21 FNEFVNDNKLTIFYDDEFQFMQKMLKFD-KDVLAIVNEKVFKGFSKDELSDL--LFFKSF 77
F + + + K T D + M + K K D DV + NEK + + L + + L + + + KK

Sbjct: 665 FEKYCSNIKNTLIRDD--MKKFRKPDISDVHILHNEKIYLEKLLNEKLNKLYIKDIEKKLD 721

Query: 78 TIHFLDREINRQTVEAFGMQV-----ITVCITHEDYLNVVYSSEVEKYLQSQGFTEHNE 132
+ H + IN + + + + QV I V + DY + S + + K + + N

Sbjct: 722 ELHGV---INKNKEDIYILQVEKQTLIKVISSVYDYTKME-SENHIFKMNITWNKMLNNV 777

Query: 133 DTSNTDETSNQATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKT 192
+ SN D + NQN + + + N + + N N + + + N + N + N

Sbjct: 778 HMSSNKDY-NNQNNQNIENNQNIENNQN-----NQNIEN-----NQNIENNQNN 820

Query: 193 VNKSSNESQNAKRQKQKGN 214
N + N + + NQN + NQN + NA

Sbjct: 821 QNNQNNQNNQNNQNNQNNQNNNA 842

Score = 33.6 bits (75), Expect = 1.4
Identities = 46/221 (20%), Positives = 89/221 (39%), Gaps = 37/221 (16%)

Query: 10 DFIKSELIKGFNEFVNDNKLTIFYDDEFQFMQKMLKFDKDVLAIVNEKVFKGFSKDELS 69
D + K E K N + + L Y + + M + K K + V K SL

Sbjct: 367 DSLKIEYNKSKTNIQQLNEQLVNYKNFIKEMEKKYK-----QLVVKNNSLFSITH 416

Query: 70 DLLFKKSFTIHLFLDREINRQTVEAFGMQVITVCITH---EDYLNVVYSSEVEKYLQSQG 126

Score = 32.8 bits (73), Expect = 2.4
Identities = 28/122 (22%), Positives = 53/122 (42%), Gaps = 2/122 (1%)

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Query: 119  EKYLQSQGFTEHNEEDTTSNTDETSNQNATSLDNSTGMTANRNAVVSLLPQSEVNID-VDNT 177
           E Y S + + + + N + +N + + DN+ N N ++ +N D ++N
Sbjct: 2838  EYYPVSTHYDNNDDINKDNIINNDDNNNDINDDNNNDINNNDDNNNDINNNDDINND 2897

Query: 178  TLRFADNNTIIDNGKTVNKSSNESNQAKRNQKQGNKAGTQFTKQYLIDNIDKAYDLRKK 237
           +N+ +NG SSN ++ N N N K N +G + + + + YD K
Sbjct: 2898  NNNDDNNDDNSNNGFVCELSSNINDDFNINILNVN-KDNFQGINKSNNFSTNLSEYNIDAYVK 2956

Query: 238  IL 239
           I+
Sbjct: 2957  IV 2958

```

Score = 32.5 bits (72), Expect = 3.1
Identities = 46/249 (18%), Positives = 101/249 (40%), Gaps = 31/249 (12%)

```

Query: 9      YDFIKSELIKKGNEFVNDNKLTFYDDEFFQFMQKMLKFDKDVLAIVNEKVFGFSLKDEL 68
            Y+++K      ++N      N      N      K      E      Q+ + K+ + + + +E      K      L+ +
Sbjct: 2150  YNYVK---VQNATNRDNKNK-----ERNLSQEIKYKINENIDLTSELEKNDMLNNYK 2200

Query: 69     SDL-----LPKKSFTIHFLDREINRQTVEAFGMQVITVCITHEDYLVNVVYSSEVEKYL 122
            ++L      ++K + I L      +      M+      +      +      N      +      E+ + L
Sbjct: 2201  NELKEKNEEIKYKLNNDIDMLSNNCKKLKESIMMEKYKIIMN-----NNIQEKDEIENL 2255

Query: 123    QSQGFTEHNEDTTSNTDETSQNQATSLDNSTGMTAN----RNAYVSLPQSE----VNIDV 174
            +++ +      +D +N      +      ++S M+ +      N      + +L +S      N+D+
Sbjct: 2256  KNK-YNNKLDDLINNYSVVDKSI VSCFEDSNIMSPSCNDILNVFNNLKSXNKXVCTNMMDI 2314

Query: 175    DNTTLRFADNNTIDNGKTVNKSSNESNQNAKRQNKQGNAGKTQFTKQYILIDNIDKAYDL 234
            N      +      ++I+N      +N      +N      N      N      N      K      YL++N+      D
Sbjct: 2315  CENEMDSI--SSINNNVNNINNVNNINNVNNINNVNNINNVNNKVIQDINNYLVNVLQLNKN 2372

Query: 235    RKKILNEFD 243
            I+ +F+
Sbjct: 2373  DNIIIIKFEN 2381

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Score = 32.1 bits (71), Expect = 4.1
Identities = 20/103 (19%), Positives = 48/103 (46%), Gaps = 2/103 (1%)

Query: 115 SSVVEKYQSQGFTEHNEDDTTSNTDETSQN--ATSLDNSTGMTANRNYAVSLPQSEVNI 172
+++ EKY EH ++ L D ++N+ N L ++ ++ + N S ++B+
Sbjct: 3264 DDDEKYSCHDDKNEHTNNDLLNDHDHNNKNNITDLEYTVVSVSHNKPSPKENEIQ 3323

Query: 173 DVDNTTLRFADNNTIDNGKTVKSSNESNQAKRNNQKGNK 215
+ + D N ++ N ++E++N + ++N + + K
Sbjct: 3324 LISIDSSNENDENDENDENDENDENDENDENDENDENDEK 3366

Score = 30.9 bits (68), Expect = 9.2
Identities = 27/118 (22%), Positives = 53/118 (44%), Gaps = 15/118 (12%)

Query: 104 THEDYLVNVVYSSSEV----EKYLQSQGFTEHNEDTTSNTDETSQNQATSLDNSTGMTANR 159
T+D LN++ + +++ E Y HN+HDPK ++ KEE QN S+D+S N
Sbjct: 3280 TNDLLNIIDHDNNKNINITDELYSTYNFVSNNHKPPSNKEII--QLNISIDSSNENDEND 3337

Query: 160 NAYVSLPQSEVNIDVDNTLRFADNNTIDNGKITVKNSSNESNQNAKRNNQKGNAGKT 217
+++ N+N+D DN ++ N +++++ + ++ N +GT
Sbjct: 3338 EN-----DENDENDEN-----DENDENDENEKDENDENDENFDNNNEG 3386

302

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>gi|585795|sp|P21538|REB1_YEAST DNA-BINDING PROTEIN REB1 (QBP)
>gi|626139|pir|[S45907 DNA-binding protein REB1 - yeast
(Saccharomyces cerevisiae) >gi|536280|emb|CAA84992|
(Z35918) ORF YBR049c [Saccharomyces cerevisiae]
>gi|559944|emb|CAA86391| (Z46260) REB1 DNA-binding
protein [Saccharomyces cerevisiae]
Length = 810

Score = 45.7 bits (106), Expect = 3e-04
Identities = 34/158 (21%), Positives = 72/158 (45%), Gaps = 14/158 (8%)

Query: 83 DREINRQTVEAFGMQVITVCITHEDYLVVYSSEVEKYLSQGFTEHNEDTTSNTDETS 142
D+ N+++VE ++ + V + H+++ +++ K+ + Q E + D N ++ S
Sbjct: 7 DKNANQESVVEAVLKVVGVLHDQNHDPQLHTKDLENKHSKKQNVESSSDQVNNNDSS 66

Query: 143 NQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID---NGKTVNKSSNE 199
N+N + D+S ++A L +E + +VD+ N +D N+ +E
Sbjct: 67 NRNEDNNDDSENISA-----LNANESSSNVDHANSNEQHNAVMDWYLRQTAHNQQDDE 119

Query: 200 SNQNAKRNNQKGNAGTQFTKQYLIDNIDKAYDLRKK 237
++N N GN F++ ++ +D D KK
Sbjct: 120 DDEN--NNNTDNGNDSNNHFSQSDIV--VDDDDDKNKK 153

>gi|172372 (M58728) DNA-binding protein [Saccharomyces cerevisiae]
Length = 809

Score = 45.7 bits (106), Expect = 3e-04
Identities = 34/158 (21%), Positives = 72/158 (45%), Gaps = 14/158 (8%)

Query: 83 DREINRQTVEAFGMQVITVCITHEDYLVVYSSEVEKYLSQGFTEHNEDTTSNTDETS 142
D+ N+++VE ++ + V + H+++ +++ K+ + Q E + D N ++ S
Sbjct: 7 DKNANQESVVEAVLKVVGVLHDQNHDPQLHTKDLENKHSKKQNVESSSDQVNNNDSS 66

Query: 143 NQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID---NGKTVNKSSNE 199
N+N + D+S ++A L +E + +VD+ N +D N+ +E
Sbjct: 67 NRNEDNNDDSENISA-----LNANESSSNVDHANSNEQHNAVMDWYLRQTAHNQQDDE 119

Query: 200 SNQNAKRNNQKGNAGTQFTKQYLIDNIDKAYDLRKK 237
++N N GN F++ ++ +D D KK
Sbjct: 120 DDEN--NNNTDNGNDSNNHFSQSDIV--VDDDDDKNKK 153

>gi|2952545 (AF051898) coronin binding protein [Dictyostelium
discoideum]
Length = 560

Score = 44.9 bits (104), Expect = 6e-04
Identities = 26/83 (31%), Positives = 39/83 (46%), Gaps = 5/83 (6%)

Query: 131 NEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNG 190
N + +N +N N+ S +NS +N N+ + P N D DN T +NNT +N
Sbjct: 404 NNNNNNNIINNNSNSNSNNNSNN-NSNNNSNRNSPNKNNNGDNDNNT---NNNTNNNN 458

Query: 191 KTVNKSSNESNQNKRNNQKGN 213
N ++N +N N N N N
Sbjct: 459 NNNNNNNNNNNNNNNNNNNNNNN 481

Score = 41.4 bits (95), Expect = 0.006
Identities = 22/88 (25%), Positives = 43/88 (48%), Gaps = 6/88 (6%)

Query: 130 HNEDTTSNTDETSNQNATSLDN---STGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNT 186
+ ++ +N++ SN N+ + +N + G AN++ + P + +N + DN +NN
Sbjct: 337 NRNNSNNNSNNNSNNNSNNNSNNRNITGNSNANKS---NSPNNNLNTNNDNKNNNSNNNNNN 393

Query: 187 IDNGKTVNKSSNESNQNKRNNQKGN 214
+N S+N +N N N N N+
Sbjct: 394 SNNNSNNGNSNNNNNNNNIINNNSNSNS 421

Score = 40.6 bits (93), Expect = 0.011
Identities = 24/101 (23%), Positives = 41/101 (39%), Gaps = 2/101 (1%)

Query: 115 SSEVEKYLSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDV 174
S+ L + ++N +N ++ N S +N+ N N S + N +

```


Query: 145 NATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVNKSSNESNQNA 204
N + NSTG T+ NI + N L +N T + T + ++ +N N+

305

Sbjct: 176 NINTNTNSTGNTSTTKKLT-----NI-ITNQILTGNNNTTNTSSTEHNNNINTNTNS 228

Query: 205 KRQKQKGNAGTQFTKQYLIDNIDKAYDL 234
N N N T + DNI+ +L

Sbjct: 229 TDNSNTNTNLTDITTTTKKWDNINTTQNL 258

Score = 41.8 bits (96), Expect = 0.005
Identities = 30/101 (29%), Positives = 43/101 (41%), Gaps = 13/101 (12%)

Query: 130 HNEDETTSTNDETSNQATSLDNTGTMANRNAYVSLPQSEVNIDV-----DNTTLRFA 182
+N DT S ++ ++ AT DN+ T T N N + N D +NT +

Sbjct: 363 NNTDTISTDNDNTDKATDNDNTDKATDNNNTDTKATDNNNTDTKATDKSNTDTKAT 422

Query: 183 DNN-----TIDNGKTVNKSSNESQNAKRQKGNAGT 217
DNN DN T K+++ +N N K N N K T

Sbjct: 423 DNNNTDTKATDNNNTNTKATDSNNTNTKATDNNNTNTKAT 463

Score = 40.6 bits (93), Expect = 0.011
Identities = 31/121 (25%), Positives = 47/121 (38%), Gaps = 31/121 (25%)

Query: 128 TEHNEDTTSTNDETSNQAT-----SLDNTGTMANRNAYVSLPQSEVN----- 171
TEHN + +NT+ T N + T ++ + +T N N + +E N

Sbjct: 171 TEHNNNINTNTNSTGNTSTTKKLTENIITNQILTGNNNTTNTSSTEHNNNINTNTNSTD 230

Query: 172 -----IDVDNTTLRFADN-----NTIDNGKTVNKSSNESQNAKRQKGNAGT 216
D+ TT ++ DN T N TV+ +N +N N K N N K

Sbjct: 231 NSNTNTNLTDITTTTKKWDNINTTQNLTTSTNTTTVSTDNNNNINTKPTDNNNTNIKS 290

Query: 217 T 217

T

Sbjct: 291 T 291

Score = 38.3 bits (87), Expect = 0.055
Identities = 28/98 (28%), Positives = 41/98 (41%), Gaps = 10/98 (10%)

Query: 128 TEHNEDTTSTNDETSNQATSLDNTGTMANRNAYVSLPQSEVNIDVD-NTTLRFADNNT 186
TEHN + +NT+ S N+ + N T +T + + N+ NTT DNN

Sbjct: 216 TEHNNNINTNTN--STDNSNTNTNLTDITTTTKKWDNINTTQNLTTSTNTTTVSTDNNN 273

Query: 187 -----IDNGKTVNKSSNESQNAKRQKGNAGT 217
DN T KS++ N K N+ + K T

Sbjct: 274 NNINTKPTDNNNTNIKSTDNYNTGTKETDNKNTDIKAT 311

Score = 37.5 bits (85), Expect = 0.094
Identities = 31/106 (29%), Positives = 45/106 (42%), Gaps = 18/106 (16%)

Query: 128 TEHNEDTTSTNDETSNQATSLDNTGTMANRNAYVSLPQSEVN-----IDVDN 176
T++N +T +T T N N AT N+T A N + ++ N D +N

Sbjct: 390 TDNNNT--DTKATDNNNTDTKATDKSNTDTKATDNNNTDTKATDNNNTNTKATDSNN 447

Query: 177 TTLRFADNN-----TIDNGKTVNKSSNESQNAKRQKGNAGT 217
T + DNN DN T K+++ +N N K N N K T

Sbjct: 448 TNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKAT 493

Score = 35.2 bits (79), Expect = 0.47
Identities = 24/109 (22%), Positives = 46/109 (42%), Gaps = 6/109 (5%)

Query: 128 TEHNEDTTSTNDETSNQATSLDNTGTMANRNAYVSLPQSEVN-----IDVDNTTLRF 181
T++N T TD + + +N+T A N + ++ N D +NT +

Sbjct: 473 TDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKA 532

Query: 182 ADNNTIDNGKTVNKSSNESQNAKRQKGNAGTQFTKQYLIDNIDK 230
DNN N + +E+ + K N++ N++ + K + +DK

Sbjct: 533 TDNNNTNQYVFANNYDETTSDDKLNKDCDENSEEKENIKSMINAYLDK 581

Score = 34.4 bits (77), Expect = 0.81
Identities = 26/126 (20%), Positives = 46/126 (35%), Gaps = 7/126 (5%)

306

Query: 99 ITVCITHEDYLNVVYSSSEVEKYLSQSGFTEHNEDTTSNTDETSNQATSLDNSTGMTAN 158
IT T+ ++ S+ V S T +++ +N T N N ++ T
Sbjct: 318 ITTDTNTNTVISTDNSKTNVISKDNSNTHITISTDNSKTNVISTDNNNTDTISTDNDNTDT 377

Query: 159 RNAYVSLPQSEVNIDVDNTTLRFADNNTID-----NGKTVNKSSNESQNAKRQK 211
+ ++ + +NT + DNN D N + N +N + K N
Sbjct: 378 KATDNDNTDTKATDNNNTDTKATDNNNTDTKATDKSNNTDTKATDNNNTDTKATDNNN 437

Query: 212 GNAKGT 217
N K T
Sbjct: 438 TTKAT 443

Score = 34.4 bits (77), Expect = 0.81
Identities = 30/100 (30%), Positives = 44/100 (44%), Gaps = 14/100 (14%)

Query: 131 NEDTTSNTDETSNQATSLDNS-TGMTANRNAY---VSLPQSEVNI---DVDNTTLRFAD 183
N + T TD T N N S DNS T + + N+ +S S+ N+ D +NT D
Sbjct: 313 NNNITITDNT-NTNVISTDNSKTNVISKDNSNTHITISTDNSKTNVISTDNNNTDTISTD 371

Query: 184 NNTIDNGKTVNKSS-----NESQNAKRQKGNAGKT 217
N+ D T N ++ N +N + K N + K T
Sbjct: 372 NDNTDTKATDNDNTDTKATDNNNTDTKATDNNNTDTKAT 411

Score = 34.4 bits (77), Expect = 0.81
Identities = 28/101 (27%), Positives = 41/101 (39%), Gaps = 15/101 (14%)

Query: 131 NEDTTSNTDETSNQATSLDNSTGMTA--NRNAYVSLPQSEVNIDV-----DNTTLRFA 182
N DT + ++ ++ AT +N+T A N N N D +NT +
Sbjct: 374 NTDTKATDNDNTDTKATDNNNTDTKATDNNNTDTKATDKSNNTDTKATDNNNTDTKAT 433

Query: 183 DNNITIDNGK-----TVNKSSNESQNAKRQKGNAGKT 217
DNN N K T K+++ +N N K N N K T
Sbjct: 434 DNNN-TTKATDSNNTNTKATDNNNTNTKATDNNNTNTKAT 473

Score = 32.5 bits (72), Expect = 3.1
Identities = 30/110 (27%), Positives = 40/110 (36%), Gaps = 23/110 (20%)

Query: 131 NEDTTSNTDETSNQATSLDNS-----TGMTANRNAYVSLPQS----EVNIDVDNTTLRF 181
N +TT N ++N S DN+ T T N N + + D NT ++
Sbjct: 251 NINTTQNLTTSTNTTTVSTDNNNNINTKPTDNNNTNIKSTDNYNTGKETDNKNTDIKA 310

Query: 182 ADNNTI-----DNGKTVNKSSNESQNAKRQKGNAGKT 217
DNN I DN KT S + SN + N K N T
Sbjct: 311 TDNNNTITDNTNTNVISTDNSKTNVISKDNSNTHITISTDNSKTNVIST 360

>gi|1429240|emb|CAA67659| (X99260) lower collar protein
[Bacteriophage B103]
Length = 293

Score = 43.8 bits (101), Expect = 0.001
Identities = 53/204 (25%), Positives = 79/204 (37%), Gaps = 42/204 (20%)

Query: 56 EKVPKG----FSLKDELSDDLFPKKSFTIHFLD----REINRQTVEAFGMQVITVCITHED 107
EK+ KG F + + D ++K F HF+ REI +T F + T I +
Sbjct: 26 EKIEKGRPKLFDFQYPIFDESIRKVFETHFIRNFYMRIGFETEGLFKFNLETWLIINMP 85

Query: 108 YLNVVYSSSEVEKY-----LQSGFTEH-----NEDTT-----SNTDETSNQNA 146
Y N ++ S E+ KY L + G ++ N DTT SNT + NA
Sbjct: 86 YFNKLFESE-LIKYDPLENTRLNNTGNKKNDTERNDNRDTTGSMDKGSNTKTSKDTNA 144

Query: 147 TSLDNSTGMTA-----NRNAYVSLPQSEVNIDVDN--TTLRFADNNTIDNGKTVNKS 196
T G T NR P S +N+ ++ TL +A + I+ T NK
Sbjct: 145 TGSSKEGDKTTGVSITDNNFNKIDSQDPSRLNLTNDGQGTLEYA--SAIEENNTNNKR 202

Query: 197 SNESQNAKRQKGNAGKTQFT 220
+ N + + GT T
Sbjct: 203 NTTGTNNVTSSAESESTGSGTSDT 226

Query= pt|110879 44AHJDORF009 Phage 44AHJD ORF |5744-6496|2 1
(250 letters)

307

>gi|2764981|emb|CAA69021.1| (Y07739) N-acetylmuramoyl-L-alanine
amidase [Staphylococcus phage Twort]
Length = 467

Score = 180 bits (452), Expect = 1e-44
Identities = 89/157 (56%), Positives = 109/157 (68%), Gaps = 8/157 (5%)

Query: 1 MKSQQAQKEWIYKHEGAGVDFGAYGFQCMDLSVAYVYIITDGKVRMWNAGDAINNDFK 60
MK+ +QA+ +I G DFDG YG+QCMDL+V Y+Y++TDGK+RMWGNAGDAINN F
Sbjct: 1 MKTLKQAESYIKSVNTGTDFDGLYGYQCMDLAVDYIYHVTGKIRMWGNAGDAINNSFG 60

Query: 61 GLATVYKNTPSFKPQLGDAVYVYNGQ---YGHICVLS---GNLDYITCLEQNWLGSGF 113
G ATVYKN P+P+P+ GDV V+T G YGHI V + G+L Y T LEQNW G G
Sbjct: 61 GTATVYKNTPAFRPKYGDVVVWTTGNFATYGHIAIVTNPDPYGLQYVTVLEQNWNGNGI 120

Query: 114 DGWEKATIRTHYDGVTHFIRPKFSGSNS-KALETSK 149
E ATIRTH Y G+THFIRP F+ +S K +T K
Sbjct: 121 YKTELATIRTHDYTGITHFIRPNFATESSVKKKDTKK 157

Score = 61.7 bits (147), Expect = 6e-09
Identities = 41/125 (32%), Positives = 57/125 (44%), Gaps = 8/125 (6%)

Query: 125 YYDGVTHFIRPKFSGSNSKALETSKVNITFGKWRNQGTYRNGTFTC-GFLPIFARV 183
YY+G T P +K + +T G W N YGTY++E+ TF C I R
Sbjct: 346 YYEGKTPV--PTVVNQKAKTKPVKQSTSG-WNVNNGTYKSESATFKCTARQGIVTRY 402

Query: 184 GSPKLSEPNQYWFQPNQYTPYNEVCLSDGYVWIGYNWQGT-YYLPVRQWNGKTGNSYSV 242
P + P Y+ VC DGYVMI + G + ++PVR W+ N+ +
Sbjct: 403 TGPFTTCCPQAGVLYGQSVTYDTVCKQDGYVWISWTTNGGQDVWMPVRTWD---KNTDIM 459

Query: 243 GIPWG 247
G WG
Sbjct: 460 GQLWG 464

>gi|113675|sp|P24556|ALYS_STAAU AUTOLYSIN
(N-ACETYLMURAMOYL-L-ALANINE AMIDASE)
>gi|79887|pir|JQ1147 N-acetylmuramoyl-L-alanine amidase
(EC 3.5.1.28) - Staphylococcus aureus >gi|153067
(M76714) peptidoglycan hydrolase [Staphylococcus aureus]
Length = 481

Score = 118 bits (292), Expect = 6e-26
Identities = 56/117 (47%), Positives = 68/117 (57%), Gaps = 1/117 (0%)

Query: 135 PKFSGSNSKALETSKVNITFGK-WKRNQYGTYYRNGTFTCGFLPIFARVGSPLSEPNG 193
P + SN + ++ V WKRN+YGTY E+ FT G PI R P LS P G
Sbjct: 365 PVATVSNESASSNTVKPVASAWKRNKYGTYYMEESARFTNGNQPIVTRKVGPFLLSCPVG 424

Query: 194 YWFQPNQYTPYNEVCLSDGYVWIGYNWQGTYYLPVRQWNGKTGNSYSVGIPIGWVFS 250
Y FQP GY Y EV L DG+VW+GY W+G RYLLP+R WNG + +G WG S
Sbjct: 425 YQFPQGGYCDYTEVMLQDGHVWVGTYWEGQRYLPPIRTWNGSAPPNQILGDLWGEIS 481

Score = 78.0 bits (189), Expect = 7e-14
Identities = 48/109 (44%), Positives = 62/109 (56%), Gaps = 6/109 (5%)

Query: 15 EGAGVDFGAYGFQCMDLSVAYVYIITDGKVRMWNAGDA-INNDFKGLATVYKNTPSFK 73
EG + D YGFQC D + A + + G + AKD N+F GLATVY+NTP F
Sbjct: 18 EGKQFNVDLWYGFQCFDYANAG-WKVLFGLLKGLGAKDIPFANNFDGLATVYQNTPDFL 76

Query: 74 PQLGDAVYVYNGQ---YGHICVLSGNLDYITCLEQNWLGSGF-DGWEK 118
Q GD+ V+ + YGH+ V+ LDY EQNWLGSG+ DG E+
Sbjct: 77 AQPQDMVVFSGSYGAGYGHVAVVIEATLDYIIIVEQNWLGGGWTGIEQ 125

>gi|1763243 (U72397) amidase [bacteriophage 80 alpha]
Length = 481

Score = 118 bits (292), Expect = 6e-26
Identities = 56/117 (47%), Positives = 68/117 (57%), Gaps = 1/117 (0%)

Query: 135 PKFSGSNSKALETSKVNITFGK-WKRNQYGTYYRNGTFTCGFLPIFARVGSPLSEPNG 193
P + SN + ++ V WKRN+YGTY E+ FT G PI R P LS P G
Sbjct: 365 PVATVSNESASSNTVKPVASAWKRNKYGTYYMEESARFTNGNQPIVTRKVGPFLLSCPVG 424

Query: 194 YWFQPNQYTPYNEVCLSDGYVWIGYNWQGTTRYLPVRQWNGKTGNSYSVGIPWGVFS 250
 Y FQP GY Y EV L DG+VW+GY W+G RYYLP+R WNG + +G WG S
 Sbjct: 425 YQFPQGGYCDYTEVMLQDGHVWVGTYWEGQRYLPRTWNGSAPPNQILGDLWGEIS 481

Score = 83.5 bits (203), Expect = 2e-15
 Identities = 50/115 (43%), Positives = 65/115 (56%), Gaps = 6/115 (5%)

Query: 9 EWYKHEGAGVDFDGYGFCMDLSVAYVYYITDGKVRMWGNAKDA-INNDFKGLATVYK 67
 EW+ EG + D YGFQC D + A + + G + AKD N+F GLATVY+
 Sbjct: 12 EWLKTSEGKQFNVDLWYGFQCFDYANAG-WKVLFGLLKGLGAKDIPFANNFDGLATVYQ 70

Query: 68 NTPSFQPLQGDVAVYTNGQ---YGHQCVLSGNLDYYTCLEQNWLGGGF-DGWEK 118
 NTP F Q GD+ V+ + YGH+ V+ LDY EQNWLGGG+ DG E+
 Sbjct: 71 NTPDFLAQPGDMVVFSGNYGAGYGHVAVIEATLDYIIVYEQNWLGGGWTDGIEQ 125

>gi|4574237|gb|AAD23962.1|AF106851_1 (AF106851) LytN [Staphylococcus aureus]
 Length = 383

Score = 84.3 bits (205), Expect = 9e-16
 Identities = 48/128 (37%), Positives = 68/128 (52%), Gaps = 7/128 (5%)

Query: 15 EGAGVDFDGYGFCMDLSVAYVYYITDGKVRMWGNAKDAINNDFKGLATVYKNTPSFKP 74
 E G DFDG+YG+QC DL Y ++ ++ +G N+F A +Y NTP+FK
 Sbjct: 252 ENRGWDFDGSYGWQCFDLVNVVYWNHLYGHGLKGYGAKDIPYANNFNSEAKIYHNTPTFKA 311

Query: 75 QLGDVAVYT---NGQYGHQCVLSGNLD---YYTCLEQNWLGGGF-DGWEKATIRTHYYD 127
 + GD+ V++ G YGH VL+G+ D + L+QNW GG+ E A H Y+
 Sbjct: 312 EPGDLVVFSGRFGGGYGHATAVLNGDYDGKLMKFQSLDQNNWNGGWRKAEVAHKVVHNYE 371

Query: 128 GVTHFIRP 135
 FIRP
 Sbjct: 372 NDMIFIRP 379

>gi|3767593|dbj|BAA33856.1| (AB015195) LytN [Staphylococcus aureus]
 Length = 383

Score = 84.3 bits (205), Expect = 9e-16
 Identities = 48/128 (37%), Positives = 68/128 (52%), Gaps = 7/128 (5%)

Query: 15 EGAGVDFDGYGFCMDLSVAYVYYITDGKVRMWGNAKDAINNDFKGLATVYKNTPSFKP 74
 E G DFDG+YG+QC DL Y ++ ++ +G N+F A +Y NTP+FK
 Sbjct: 252 ENRGWDFDGSYGWQCFDLVNVVYWNHLYGHGLKGYGAKDIPYANNFNSEAKIYHNTPTFKA 311

Query: 75 QLGDVAVYT---NGQYGHQCVLSGNLD---YYTCLEQNWLGGGF-DGWEKATIRTHYYD 127
 + GD+ V++ G YGH VL+G+ D + L+QNW GG+ E A H Y+
 Sbjct: 312 EPGDLVVFSGRFGGGYGHATAVLNGDYDGKLMKFQSLDQNNWNGGWRKAEVAHKVVHNYE 371

Query: 128 GVTHFIRP 135
 FIRP
 Sbjct: 372 NDMIFIRP 379

>gi|2764983|emb|CAA69022.1| (Y07740) cell wall hydrolase Ply187
 [Staphylococcus phage 187]
 Length = 628

Score = 76.9 bits (186), Expect = 2e-13
 Identities = 50/144 (34%), Positives = 68/144 (46%), Gaps = 18/144 (12%)

Query: 5 QQAKEWIYKHEGAGVDFDGYGFCMDLSVAYVYYITDGKVRMW-----GNAKDAINNDF 59
 +Q +W G+GVD DG YG QC DL Y++ R W GNA+D +
 Sbjct: 12 KQVVDWAINLIGSGVDVGGYGRQCWDLN-NYIFN-----RYWNFKTPGNARDMAWYRY 64

Query: 60 KGLATVYKNTPSFKPQLGDVAVYTNGQY-----GHQCVLS-GNLDYYTCLEQNWLGGGF 113
 V++NT F P+ GD+AV+T G Y GH V+ Y+ ++QNW
 Sbjct: 65 PEGFKVFRNTSDFVPKPGDIAVWTGGYNWNTWGTGIVVGPSTKSYFYSVDQNNWNSNS 124

Query: 114 DGWEKATIRTHYYDGVTHFIRPKF 137
 A H Y GVTHF+RP +
 Sbjct: 125 YVGSAAKIKHSYFGVTHFVRPAY 148

>gi|3287732|sp|O05156|ALE1_STACP GLYCYL-GLYCINE ENDOPEPTIDASE ALE-1
 PRECURSOR >gi|1890068|dbj|BAA13069| (D86328) ALE-1
 [Staphylococcus capitis]
 Length = 362

Score = 73.4 bits (177), Expect = 2e-12
 Identities = 47/117 (40%), Positives = 61/117 (51%), Gaps = 10/117 (8%)

Query: 132 FIRPKFSGSNSKALETSKVNTFGKWKRNQYGYTYRNENGTFTCGFLPIFARVGSPLKSEP 191
 F++ GSNS TS N G +K N+YGT Y++E+ +FT I R+ P S P
 Sbjct: 252 FLKSAGYGSNS----TSSNNNG-YKTNKYGTLYKSESASFTAN-TDIITRLTGPFPSMP 305

Query: 192 NGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYLPPVRQWNGKTGNSYSVGIPWG 247
 + Y+EV DG+VW+GYN G R YLPVR WN TG +G WG
 Sbjct: 306 QSGVLRKGLTIKYDEVKQDGHVWVGYNNSGKRVYLPVRTWNESTG---ELGPLWG 359

>gi|79926|pir||A25881 lysostaphin precursor - Staphylococcus
 simulans >gi|153047 (M15686) lysostaphin (ttg start
 codon) [Staphylococcus simulans]
 Length = 389

Score = 69.5 bits (167), Expect = 3e-11
 Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)

Query: 131 HFIRPKFSGSNSKALETSKVNTFGK-----WKRNQYGYTYRNENGTFTCG 175
 HF R S SNS A + K +GK WK N+YGT Y++E+ +FT
 Sbjct: 258 HFQRMVNSFSNSTAQDPMFPLKSAGYKAGGTVTPTPTNGWKTNKYGTLYKSESASFTPN 317

Query: 176 FLPIFARVGSPLKSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYLPPVRQWNG 234
 I R P S P + Y+EV DG+VW+GY G R YLPVR WN
 Sbjct: 318 -TDIITRTTGPFPSMPQSGVLKAGQTIHYDEVKQDGHVWVGTYGNSGQRIYLPVRTWNK 376

Query: 235 KTGNSYSVGIPWG 247
 T ++G+ WG
 Sbjct: 377 STN---TLGVLWG 386

>gi|126496|sp|P10548|LSTP_STAST LYSOSTAPHIN PRECURSOR
 (GLYCYL-GLYCINE ENDOPEPTIDASE) >gi|79927|pir||S01079
 lysostaphin precursor - Staphylococcus simulans bv.
 staphyloolyticus >gi|581744|emb|CAA29494| (X06121)
 lysostaphin (AA 1-480) [Staphylococcus simulans bv.
 staphyloolyticus]
 Length = 480

Score = 69.5 bits (167), Expect = 3e-11
 Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)

Query: 131 HFIRPKFSGSNSKALETSKVNTFGK-----WKRNQYGYTYRNENGTFTCG 175
 HF R S SNS A + K +GK WK N+YGT Y++E+ +FT
 Sbjct: 349 HFQRMVNSFSNSTAQDPMFPLKSAGYKAGGTVTPTPTNGWKTNKYGTLYKSESASFTPN 408

Query: 176 FLPIFARVGSPLKSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYLPPVRQWNG 234
 I R P S P + Y+EV DG+VW+GY G R YLPVR WN
 Sbjct: 409 -TDIITRTTGPFPSMPQSGVLKAGQTIHYDEVKQDGHVWVGTYGNSGQRIYLPVRTWNK 467

Query: 235 KTGNSYSVGIPWG 247
 T ++G+ WG
 Sbjct: 468 STN---TLGVLWG 477

>gi|3287967|sp|P10547|LSTP_STASI LYSOSTAPHIN PRECURSOR
 (GLYCYL-GLYCINE ENDOPEPTIDASE) >gi|2072411 (U66883)
 lysostaphin [Staphylococcus simulans]
 Length = 493

Score = 69.5 bits (167), Expect = 3e-11
 Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)

Query: 131 HFIRPKFSGSNSKALETSKVNTFGK-----WKRNQYGYTYRNENGTFTCG 175
 HF R S SNS A + K +GK WK N+YGT Y++E+ +FT
 Sbjct: 362 HFQRMVNSFSNSTAQDPMFPLKSAGYKAGGTVTPTPTNGWKTNKYGTLYKSESASFTPN 421

Query: 176 FLPIFARVGSPLKSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYLPPVRQWNG 234

310

I R P S P + Y+EV DG+VW+GY G R YLPVR WN
Sbjct: 422 -TDIIITRTTGPFRSMPSQGV LKAGQTIHYDEVKQDGHVWVG YTGNSGQRIYLPVRTWNK 480

Query: 235 KTGNSYSVSGIPWG 247

T ++G+ WG
Sbjct: 481 STN---TLGV LMG 490

>gi|3341932|dbj|BAA31898.1| (AB009866) amidase (peptidoglycan
hydrolase) [bacteriophage phi PVL]
Length = 484

Score = 68.3 bits (164), Expect = 6e-11
Identities = 52/150 (34%), Positives = 71/150 (46%), Gaps = 17/150 (11%)

Query: 3 SQQQAKEWIYKHGAGVDFD GAYGFQCMDLSVAVVYITDGKVRMWGNKADAINNDFKGL 62
++ QA++W G + D YGFQC D + + + I G+ R+ G I D K

Sbjct: 4 TKNQAEKWFDNSLGKQFNPD L FYGFQCYDYASMF-FMIATGE-RLQGLYAYNIPFDNKAR 61

Query: 63 ATVY----KNTPSFKPQLGDVA VYTN---GQYGHICVLSGNLDYYTCLEQNWLGGGF-- 113

Y KN SF PQ D+ V+ + G GH++ V S NL+ +T QNW G G+
Sbjct: 62 IEKYGGIIRKND SFLPQKLDIVVFP SKYGGGAGHVEIVESANLNTFTSFGQNWNGKGWTN 121

Query: 114 ----DGW--EKATIRTHYDGVTHFIRPKF 137

GW E T HYYD +FIR F
Sbjct: 122 GVAQPGWGPETVTRHVHYDDPMYFIRLNF 151

Query= pt|110882 44AHJDORF012 Phage 44AHJD ORF |8391-8813|3 1
(140 letters)

>gi|140528|sp|P24811|YQXH_BACSU HYPOTHETICAL 15.7 KD PROTEIN IN
SPOIIIC-CWLA INTERGENIC REGION (ORF2)
>gi|322189|pir|B44816 orf2 5' of autolytic amidase -
Bacillus subtilis >gi|142801 (M59232) open reading frame
2 [Bacillus subtilis] >gi|1217874|dbj|BAA06959| (D32216)
ORF121 [Bacillus subtilis] >gi|1303767|dbj|BAA12423|
(D84432) YqdD [Bacillus subtilis]
>gi|2635036|emb|CAB14532| (Z99117) alternate gene name:
yqdD; similar to holin [Bacillus subtilis]
Length = 140

Score = 80.4 bits (195), Expect = 6e-15
Identities = 45/130 (34%), Positives = 67/130 (50%), Gaps = 3/130 (2%)

Query: 4 VKFRPTDSEAFHMFYAGDLKLLYFLFVLMFVDIITGSKAIKNNLWSKKSMRGFSKKX 63

+ F D ++P G +K L L VL +D++TG+ KA K L S+ + G+ +K
Sbjct: 8 INFETLDLARVYLF---GGVKYLDLLVLVLSIIDVLTGVKAWKFKLRSRS AWFYVRKL 64

Query: 64 XXXXXXXXXXXXXXXXXXXXGGLLMITIFYIANEGLSIVENCAEMDVLVPEQIKDKLRVI 123

G L T+ +YIANEGLSI EN A++ V +P I D+L+ I
Sbjct: 65 LNFFAVILANVIDTVLNLNGVLTFTGTVLFYIANEGLSITENLAQIGVKIPSSITDRLQTI 124

Query: 124 KNDTEKSDNN 133

+N+ E+S NN
Sbjct: 125 ENEKEQSKNN 134

>gi|4126631|dbj|BAA36651.1| (AB016282) ORF45 [bacteriophage phi-105]
Length = 135

Score = 76.1 bits (184), Expect = 1e-13
Identities = 44/115 (38%), Positives = 61/115 (52%), Gaps = 4/115 (3%)

Query: 21 GDLKLLYFLFVLMFVDIITGSKAIKNNLWSKKSMRGFSKKXXXXXXXXXXXXXXXXXXXX 80

G++K L + VL +DIITG+ KA K L S+ + G+ +K
Sbjct: 17 GEVKYLDLMLVLNIIIDITGVKAWKFKELRSRS AWFYVRKMLSFVVIVANAIDTIMD 76

Query: 81 XKGGLLMITIFYIANEGLSIVENCAEMDVLVPEQIKDKLRVIKND----TEKSD 131

G L T+ +YIANEGLSI EN A++ V +P I D+L VI++D TEK D
Sbjct: 77 LNGVLT FATVLFYIANEGLSITENLAQIGVKIPAVITDRLHVIESDNDQKTEKDD 131

>gi|141088|sp|P26835|YNGD_CLOPE HYPOTHETICAL 14.9 KD PROTEIN IN NAGH
3'REGION (ORFD) >gi|1075967|pir||S43905 hypothetical
protein D - Clostridium perfringens >gi|455154 (M81878)

311

ORF D [Clostridium perfringens]
Length = 132

Score = 60.9 bits (145), Expect = 4e-09
Identities = 38/127 (29%), Positives = 63/127 (48%), Gaps = 3/127 (2%)

Query: 1 MNEVKFRPTDSEAFHMFYI-AGDLKLLYFLFVLMFVDIITGSKAIKNNNLWSKKSMRGF 59
+N +K+ +I+ A D+ L+ L V +F+D +TG+ K K+ L S +RG
Sbjct: 5 INYIKWGIIVSLGTLFTWIFGAWDIPLITLL-VFIFLDYLTGVKCKSKELCSNIGLRGI 63

Query: 60 SKXXXXXXXXXXXXXXXXXXXXGGLLMITI-FYYIANEGLSIVENCAEMDVLVPEQIKD 118
+KK + I ++YI NEG+SI+ENCA + V +PE++K
Sbjct: 64 TKKGLILVLLVAVMLDRLLDNGTWMFRTLIAFYIMNEGISILENCAALGVPIPEKLKQ 123

Query: 119 KLRVIKN 125
L+ + N
Sbjct: 124 ALKQLNN 130

>gi|2293160 (AF008220) YtkC [Bacillus subtilis]
>gi|2635548|emb|CAB15042| (Z99119) similar to autolytic
amidase [Bacillus subtilis]
Length = 134

Score = 36.4 bits (82), Expect = 0.099
Identities = 25/109 (22%), Positives = 41/109 (36%)

Query: 17 FIYAGDLKLLYFLFVLMFVDIITGSKAIKNNNLWSKKSMRGFSKXXXXXXXXXXXXXXXXX 76
F + G L LM ++ I+ K + L KK KK
Sbjct: 20 FFFGGFQYSFLILLSLMAIEPISTTLKETIIHKLSFKKVFARLVKKLVTLALISVCHFFD 79

Query: 77 XXXXXKGGLLMITIFYIYIANEGLSIVENCAEMDVLVPEQIKDKLRVIKN 125
+G + + I +YI E + IV + + + VP+ + D L +KN
Sbjct: 80 QLLNTQGSIRDLAIMFYILYESVQIVVTASSLGIPVQMLVDLLETLEKN 128

>gi|1181973|emb|CAA87743.1| (Z47794) holin protein [Bacteriophage
CP-1]
Length = 134

Score = 31.3 bits (69), Expect = 3.3
Identities = 27/88 (30%), Positives = 36/88 (40%), Gaps = 5/88 (5%)

Query: 29 LFLVLMFVDIITGSKAIKNNNLWSKKSMRGFSKXXXXXXXXXXXXXXXXXXXXK--GGLL 86
LF L+ D ITG KA K S ++G K G +L
Sbjct: 18 LFLALILFDITGFLKAWKWKVTDSTGLKGVIKHTLTFIFYFVAVFLTYIHAMAVGQIL 77

Query: 87 MITIFYIYIANEGLSIVENCAEMDVLVPE 114
++ I Y A LSI+EN A M V +P+
Sbjct: 78 LVIINLYYA---LSIMENLAVMGVFIPK 102

Table 21

Phage 182 complete genome sequence. 17833 nucleotides.

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1      tagaatattg tcataaaaca caaacataat aatgcatatt attgtttaca aatatgtaat ttcgtgatat
71     aatatatttg taagttaaag gaggtgacaa aagaacaaat cataaatgct ttagaatttg caaaaactat
141    tggaggaaaa ataattgaat attcactaca acaaatagat gaaattaaat caacaatttt cagaattaga
211    ttaaaaaggc atgaactaga ggaattgggt gacgaagtaa acgatattgc taaagatccg gaggaagat
281    atctttttac gttttattac acagaagaag aacgtttggt tgaattccoc tctgcaagat taatagatta
351    ttacaacgaa aagatcacaa atctgaaatc ggaaatcata tcactcgaaa aaagattaca aaaactagta
421    aaataattac acaaaaagct ttacaaatat aacacatcat gttatactaa aagagtagta agggaaacgga
491    aaatacctta cttcacacct caatcattct tatcaaaaata caaaaggagg gaaaataatg ggtcgaaaac
561    taatgcaacy aaacgtaaca tcaactaaag tagaattctc agaagtatc gtacaagatg gagcgccaac
631    aattgtacca tgcgaaccag ttgtcttaac aggaaaactt tcagaagaaa aagctttatc agcgatcaaa
701    cgtaaaaacc ctgataaaaa cgtagtgtga acaaatgttt cacatgaaac agcgctttac acaatgccag
771    tcgataaatt tatcgagtta gcagacaaat caacacaagc taataaaaaa caaaactaaa acaaaacaga
841    ggagattata atcatggaaa tcgtaaaaag cacatttgac acacaaacac cagaaggaat gttacaagta
911    ttcaatgccca caaacggggc ttcaattccg ttacgtaacg caattggcga agtactagaa ttgaaagata
981    ttctagttta ctcagacgaa gtttctgggt ttggtggagc cgaacctatc caagcagaac tagtcgcttt
1051   cttcacagaa gatggtaaaa cttatcgggg tgtatcagca gtagcaacaa aatcagctaa aaaccttaatt
1121   gatattgata ctgctaaccg tgacatcaaa ccaaaaattt cttttgtcga aggaaaaatc aacgggtggc
1191   aaaaatttgt aaatctacaa gtggtttcac tgtagcataa aaatacagga atctagtaag ccacttagcg
1261   aatctcgcta ggtggttttt attatgtttc tacattgagg tgtgtagaat tgaccgtaag aatatcaaa
1331   aatgatagag ccaagttaga gaaaatctac ggtaaatcta acaaaagctc taaaaaatat aatcgtttaa
1401   gacaaaaagg agttgaggaa aggcacactc caactgttcc aacatcaaa gaaaagacta ttgactacgt
1471   aaaaatcaaca aatatgagtc gtatgtattt taacaagatg ttgacgagat tggtagattt tgcacaacct
1541   tacaacgaga attacatttt tgagatcaac aagcgaaatg ttgcaatctc aagagcgcaa atcaagaag
1611   cgcaaattaa aacagagcaa gctcaaaaag cgaaagaaga acactacaaa gagcttaaca aagttaga
1681   taagaagccc acagaaaaca caattgtcac accaactatt ttaacagagt taggtgctga cttacctttt
1751   caagcaatac cagattttta tattgacgct ttcacttttc cagaaggagt tcagttctat ttagaaaaata
1821   taggaaaaaca agacgaacaa tattttgacg aaagagacca actttattac gacaatttca gacaagcgat
1891   gtttactatt ttcaattcag acgtgacgca tattgttcgt ttacttgact caatggggct tgatctattt
1961   atgaaaaaat atgttagtaa cttcttagac atgaacctg actacattta tgacgaagca gaagtacaac
2031   agaaaaaaga acaagtttac agtaagattg caaaagtgat cgagctgtaa acaggtggag aagtcctctc
2101   atataacccc acgaagaaca tcacaattaa ttcagaacaa ggagaagaat tatgattaa gaaatatact
2171   cgcaactttga aacaacaact gatctcaacg attgtcgtgt atggtcgtgg ggcgtatcgg atatagacaa
2241   cgttgacaat atgacgctcg gtttagaatt cgattctttt tttgagtggt gtaaaatgca agcgagcaca
2311   gacattttat tccacaacga aaaatttgac ggagagttta tgcctttcat gttattcaaa aatggtttca
2381   aatgggttaa agaagcaaaa gaagatcgaa cattctccac actcatatca aatatgggtc aatggtagc
2451   tttggaaatt tgttgggaag ttaattacac aacaacaaaa tcaggtaaaa cgaaaaaaga gaaatctcga
2521   acaataattt atgatagcct taaaaaatat cttttccag tgaacaaaat tgcagaagct ttaattttc
2591   ctataaaaaa aggcgaataa gattatacaa aagaagagac tattggttac aaaccaacaa aagatgaatg
2661   ggagtattta aagaacgaca ttcagattat ggcgatggca ttaaaaattc aattcgatca aggactaact
2731   cgaatgacta gaggaagcga cgtcttaggc gattacaaag attggtctaa agctacacat ggaatatcaa
2801   ctttcaaaac atggtttcct atttgtctt tagggtttga taaagactta cgtaaaagcat acaaaaggcg
2871   cttcacttgg gtaaacaaag tttttcaagg gaaagaaata ggtgacggca ttgtcttga tgtcaactct
2941   ttgtatccct ctcaaatgta cgtaagacct ttaccatag gaaacacctt attctacgaa ggagaataca
3011   aaccgaacaa cgactatccg ctgtacattc agttacattc aaaaatcaaa agtaagatc cggtttaaag agggttatat
3081   tccaaccatt caagttaagc aaagttcatt attcattcaa aacgaatatc ttgaatcaag tgaatacaag
3151   ttaggagttg acgaattaat cgactttact cttacaaatg ttgacctaga attatttttt gaacactacg
3221   atattttaga gatacattac acttacggat atatgttcaa agcttcttgt gatattgtca aaggctggat
3291   cgataaatgg atcgaaagta agaaccaccac cgaaggggct agaaaagcta acgccaaggg tatgttaaat
3361   agcttgtatg gaaagtctcg aacaaacccg gacattacag gaaaagtgc ttacatgggc gaggaacgca
3431   ttgttcgatt gacactagga gaagaagaat taagagatcc tgtttatgtt ccgcttggta gttttgtgac
3501   ggcttggggt agatatacta ccattacaac cgctcaaaaa tgttttgatc gcattattta ttgtgtatac
3571   gatagcattc atctagtagg aacagaagtt ccagaagcaa tcgatcactt ggttgatcct aaaaaacttg
3641   gttattgggg gcatgaaagc acatttcaac gagcaaaatt cattcggcag aaaacatcag tagaagaaat
3711   tgatggcgaa ttaaatgtaa agtgtgctgg tatgccagat cgaataaaa gattgttaac ttttgacaat
3781   tttgaagttg gtttttcaag ctatggaaa gttgtaccta aaagaacaca aggtggcggt gttattagtag
3851   acacaatggt tacaatcaaa taaggaggac taataatgga actatataaa gcaatgttta tctgactgta
3921   tgaagggtact attgacggtt acgatactga acactatgta gatatttttt tacatgactt tgaagaaata
3991   tatggaaaag aaacacgtga aattgaaagca gtaaccattag taaaaacagg aaatttaaaa aataaatta
4061   tttacatcct ttgcaagta tggtaaaaata ttcttgtgat agttgacaag agtcaaat tggcgagattg
4131   ggcgaatgta cacgtgaaat atcgtgcgct cccgttaagt tatggacaca taaacgtttt gaccgtcaac
4201   caatcgcaaa aaccttttag gtagtagcct taaatgtggc tactcttttt tgtgtttcag agattatgt
4271   ttacgtgtaa acagttttta tggtaataata gaatcaaaag gaggtggaga ttatggaaat taaagaactt
4341   gaatcaattt taaatggtat tcttgaagt gtcacagacg gtgaagcaag atcaaaagatt gtgaactc
4411   ttgaagcatt gcgagaagac tccggagcaa caactgaagc tttgacatca gcaaatagca cacttgaaaa
4481   gttaaagaaa gataacgaag cgttgggttat ttcaactca aaattgttcc gagaacgagc gatcgtagaa
4551   ccagcagaaa ataacgaacc agaaacagac cagaatatta cactagacga tttaggaatt taaggaggaa
4621   aaaacatggc tgacaaaatc acagaacaag atgtttctcg tgccacaaat gtagaacac cagtacaatt
4691   aatgactgct atttataata gttcatcatc tctttttcag gcgaacgtac ctatgcaaaa tgcagataac

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4761 atcgaagcgg ttggtgcagg gatcacacgt ttagacgtag taaaaaacga atttatttca acttttagttg
4831 accgtattgg taaagttagt atccgataca aatcttggcg taaccctttg aaaaagttta aaaaaggaaa
4901 catgccttta ggtcgaacga ttgaagaaat ttttgttgac attgcacagg aacataagtt caaccctgac
4971 gagtctgtta caggggtatt taaacaggaa gtccccgatg taaaaacatt gtccacgaa attaactcgtg
5041 aaggttacta caaacaaacg atccaagaag catggttaga aaaagcattt acttcattggg ataatttcaa
5111 tagtttcgtt gctggtgtaa tgaacgcttt atacacaggt gacgaagtaa gcgaatttga atacacgaaa
5181 ttattaatag caaactacca agaaaaagag ctattcaaaag agatcgaaat tggcgaaatt actgaatcaa
5251 atgcaaaaaga atttatccgt aagatcaaat caacctctaa caaattagaa tttatgagtt ccgcttacaa
5321 cgctcaagga gttaaaacat ctacctcaaa atctgatcaa tacgttatta ttgacgccga cacagacgca
5391 accattgacg ttgacgtttt agcagcgcca ttcaatatga gtaaaactga cttttagtaga caaaaaatcg
5461 ttattgatga gtttcctaaa aaagaaggcg aagaatcgtc aaatattgtg gcagttattg tagatagtga
5531 atggtttatg atctacgaca aattgtacaa aacaacaagt ctatacaacc ctgaagggtt atattggaat
5601 tattggttgc accaccacca actatattct acttctcaat tcgggaacgc tgttgccttt gttaaatcag
5671 caacaaaacc tgtcacaata gtctgtttcg caagtgcac aactagtggt gttaaaaggat catctaaaag
5741 tatcgatttg acatttcacac cagtagaagc aacaaccaa caaggagaag ttgtttcatc agcaccagca
5811 ttggttaagg caaccgtaaa acaaacagca ggtaaagcga ctgccgtaac cgtagaaggc ttagaagtcg
5881 gtcaatcatt agtaacattc acagctatcg gaggtcaaca agcaacggtt cttgttaccg tcttctctga
5951 ctaaggagga caattatggc aagaagggtat acaaatgtaa aattgttggc taacgtgcct tttgatacaa
6021 cctatacaca cacaagatgg tttaaaactc aacaggaaca ggaatcgta tttaatcgtt ttctgttct
6091 taacgagaat agagattggt cttatcaaaag ggatacacaa ctccggggag ttttttagagt agataaacac
6161 aaagacgcct tatatgcttg taactatctc atctttaaia acgaagaac ttatcctagt aaatggcagt
6231 atgcctttgt tactgatatt gaataaaga atgacaacac aagtttcgtt accttgaaa ttgatgtttt
6301 acaaaactat cgtttcgata ttggtatagc agaaagtttc attgcaaaag aacaccctca accttattat
6371 tcgaatggaa tacctttcat taatacaatt gaagagtcgc ttgattacgg tagagaatac acaacaacaa
6441 atgtaacaac ttttcaccc aacgatggag tcaattttct tgttattcta acaagtgaag caatgccagt
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7001 atgactttaa gacctgaata tcttacaggt ggtaaattga gtgtatatgt aaaaggttcg tttaggaattt
7071 ctaataaagt gatgatcgag ccgattgatt atgatgtaag taactcaacc attattacca atttaagtga
7141 caagatgtta atcgataatg atcctaacga tgtaggagtt aatctgact atgcttctgc attcattgcaa
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Table 22

Phage 182 ORFs list

nb	Name	Frame	Position	Size (a.a.)	Key words
1	182ORF001	2	5966..7780	604	Tail protein;
2	182ORF002	1	2152..3873	573	DNA polymerase;
3	182ORF003	1	11305..12639	444	
4	182ORF004	3	4626..5954	442	Major head protein;
5	182ORF005	3	12651..13700	349	Glycyl-Glycine endopeptidase; Lysostaphin precursor;
6	182ORF006	1	14995..16026	343	Encapsidation protein; ATG/GTP-binding site motif A;
7	182ORF007	1	7795..8775	326	Upper collar protein;
8	182ORF008	2	14105..14983	292	Lysozyme; Muramidase;
9	182ORF010	2	1310..2155	281	Terminal protein;
10	182ORF009	2	8765..9601	278	Lower collar protein;
11	182ORF011	1	9607..10158	183	Pre-neck appendage protein;
12	182ORF012	3	10872..11294	140	
13	182ORF013	1	10456..10860	134	
14	182ORF014	3	13716..14108	130	Lysis protein;
15	182ORF015	2	854..1225	123	Early protein;
16	182ORF018	-2	16429..16737	102	
17	182ORF020	3	10158..10454	98	Leucine-zipper motif;
18	182ORF019	3	4323..4613	96	Head protein;
19	182ORF016	-3	16749..17033	94	
20	182ORF022	1	12868..13149	93	
21	182ORF023	-2	11914..12189	91	
22	182ORF017	1	154..426	90	
23	182ORF024	3	6174..6446	90	
24	182ORF025	2	548..814	88	Early protein;
25	182ORF026	-3	12999..13259	86	
26	182ORF027	-1	14642..14896	84	
27	182ORF028	3	14430..14672	80	
28	182ORF021	-3	17106..17339	77	
29	182ORF030	-1	16199..16429	76	
30	182ORF031	-3	8379..8603	74	
31	182ORF032	-1	11195..11413	72	
32	182ORF033	-1	4727..4942	71	
33	182ORF034	-1	5951..6160	69	
34	182ORF029	-3	17412..17606	64	
35	182ORF035	-3	15570..15758	62	
36	182ORF036	-3	2127..2315	62	
37	182ORF037	-1	12095..12280	61	
38	182ORF038	3	14769..14951	60	
39	182ORF039	2	9992..10171	59	
40	182ORF040	-3	16029..16202	57	
41	182ORF041	1	3886..4056	56	Early protein;
42	182ORF042	-3	10671..10832	53	
43	182ORF043	-3	10491..10652	53	
44	182ORF044	-1	6299..6457	52	
45	182ORF045	-2	6571..6729	52	
46	182ORF046	2	2372..2527	51	
47	182ORF047	-2	13201..13353	50	
48	182ORF048	-3	3243..3395	50	
49	182ORF049	3	1578..1724	48	
50	182ORF050	2	8012..8155	47	
51	182ORF051	3	9390..9530	46	
52	182ORF052	1	4096..4233	45	
53	182ORF053	2	15656..15793	45	
54	182ORF054	-2	8002..8136	44	
55	182ORF055	2	8324..8455	43	
56	182ORF056	3	6549..6680	43	
57	182ORF057	-3	8133..8264	43	
58	182ORF058	-1	5048..5176	42	
59	182ORF059	-2	15748..15876	42	
60	182ORF060	-3	15276..15404	42	
61	182ORF061	-3	1974..2102	42	
62	182ORF062	-2	1867..1992	41	
63	182ORF063	-3	14181..14306	41	
64	182ORF064	-2	7234..7356	40	

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65	182ORF065	-2	3460..3582	40
66	182ORF066	1	4234..4353	39
67	182ORF067	-1	13763..13882	39
68	182ORF068	-1	7148..7267	39
69	182ORF069	-3	4908..5027	39
70	182ORF070	-3	912..1031	39
71	182ORF071	2	11741..11857	38
72	182ORF072	-3	11610..11723	37
73	182ORF073	-3	2763..2876	37
74	182ORF074	-1	8813..8923	36
75	182ORF075	-3	7353..7463	36
76	182ORF076	-3	2316..2426	36
77	182ORF077	2	11858..11965	35
78	182ORF078	-2	7564..7671	35
79	182ORF079	-2	7381..7488	35
80	182ORF080	-2	4372..4473	33

Table 23

Predicted amino acid sequences of ORFs from phage 182

182ORF001

5966 atggcaagaaggtatacaaatgtaaaattgttggttaacgtgccttttgataaacctatcacacacacaagatgggtttaaact
 1 M A R R Y T N V K L L A N V P F D N T Y T H T R W F K T
 6050 caacaggaaacaggaatcgactttaattcggtttcctgttcttaacgagaatagagattgttcttatcaaaaggatcacacaactc
 29 Q Q E Q E S Y F N S F P V L N E N R D C S Y Q R D T Q L
 6134 gggggaggttttagagtagataaaacacaaagacgcttatatgcttgtaactatctcatctttaaaccgaagaaacttatcct
 57 G G V F R V D K H K D A L Y A C N Y L I F K N E E T Y P
 6218 agtaaatggcagtatgcctttgttactgatatgaatataagaatgacaacacaagtttcgttacctttgaaattgatgtttta
 85 S K W Q Y A F V T D I E Y K N D N T S F V T F E I D V L
 6302 caaacttatcggttcgatattggtatacagagaagtttcattgcaaaagaacacccctcaactttattattcgaaatgaatacct
 113 Q T Y R F D I G I R E S F I A K E H P Q L Y Y S N G I P
 6386 ttcatataacaattgaagagtcgcttgattacggttagagaatacacacaacaaatgtaacaacttttcatcctaacgatgga
 141 F I N T I E E S L D Y G R E Y T T T N V T T F H P N D G
 6470 gtcaattttctgttattcttaacaagtgaaagcaggttgagataaggaagataaatcaggaggatcaatagtaggtggc
 169 V N F L V I L T S E A M P V G D K E D K S G G S I V G G
 6554 ccatctccttttcttattatcttctctatcaattcaagtgagggtatatacaaccaaattggggcagggaatgctaatttt
 197 P S P S Y Y L L P I N S S G E V Y K P N G A N F
 6638 ggagagtagcatggcgtttcttaacaacgaagaaccccttttaataagatagtcgggagtgatgtaacgtcgatatacaggata
 225 G E Y M A F L T T K E P F L N K I V G M Y V T S Y T G I
 6722 ccattcttggtgacacgcgaacaaacggtaaggtataatgcaggaggttcttataagatcatgcttccaacctacgctagt
 253 P F I V D H A N K T V R Y N A G G S Y K I M L P T Y A S
 6806 gatccaacaggaacaatgaaacattcgctttcttctgtgtaaaagaagcaagaacattcgatcctaaagaattgatcttga
 281 D P T G T M K T F A F P C V K E A R T F V P K R I D L V
 6890 gggaacgtgtataactacttttagagaagcttttcggttaaatgttaaggaatcaaaactatttatgatcctcattgtttaata
 309 G N V Y N Y F R E A F P F N V K E S K L F M Y P Y C L I
 6974 gaaattacagatacaaaaggacatgtaattgactttaagacctgaatatcttacaggtggtaaaattgagtgatataaggaagt
 337 E I T D T K G H V M T L R P E Y L T G G K L S V Y V K G
 7058 tcgtaggaatttctaataaagtgatgatcgagcggattgattgatgtaagtaactcaaccattattaccaatttaagtga
 365 S L G I S N K V M I E P I D Y D V S N S T I I T N L S D
 7142 aagatgttaatcgataatgatcctaacgatgtaggagtaaaatctgactatgcttctgcatcattgcaaggaacaaaaactcc
 393 K M L I D N D P N D V G V K S D Y A S A F M Q G N K N S
 7226 ttgattgctcaagagcaaaacattcgcaatacttcagacatggtatgggaacagtgcaatgagtagcaggaggagcgatcttt
 421 L I A Q E Q N I R N T F R H G M G N S A M S T T G G A I F
 7310 tcagccttagcaagtaacaacccctttgttggttactaacaatcatgggagcaggacaacaagtaacaacattatgttctgaa
 449 S A L A S N N P F V G L T N I M G A G Q Q V N N Y V S E
 7394 aaagaaaacggtttgaacctcttggcagggtaaagtgagatcgataatccagataatgtaacacagcttggatcaaac
 477 K E N G L N L A G K V A D I E N I P D N V T Q L G S N
 7478 ttatctttcacacaggaactttcaaaactattatcaattcgcttcaaaacaaatataatagtagtatgcaacaagacttgat
 505 L S F T T G N F Q N Y Y Q L R F K Q I K Y E Y A T R L D
 7562 cgttacttctcaatgtatggcacaagagcagtagtacaccaacttacaacaagaaagcatggaatttctattaaa
 533 R Y F S M Y G T K S N R V A T P N L Q T R K A W N P I K
 7646 ttaaaagaaccaaatattgttaggcacaatgagtagtattataacacgtgtgaacaacaaatttttagtcaggcggttacgctt
 561 L K E P N I V G T M S N D V L T R V K Q I F S A G V T L
 7730 tggcatacgaatgatgttttgaattataccaagacaacggagatgtatag 7780
 589 W H T N D V L N Y N Q D N G D V *

182ORF002

2152 atgattaagaaatatactggcgactttgaacaacaactgatctcaacgatgtcgtgtatggctcgtggggcgatgcatata
 1 M I K K Y T G D F E T T T D L N D C R V W S W G V C D I
 2236 gacaacgttgacaatatgacgttcgggttagaatacgattcttttttgagtggtgtaaaatgcaaggcagcacagacatttat
 29 D N V D N M T F G L E I D S F F E W C K M Q G S T D I Y
 2320 ttccacaacgaaaaatttgacggagaggttatgctttcatggttattcaaaatgggttcaaatgggtgtaaaagaagcaaaagaa
 57 F H N E K F D G E F M L S W L F K N G F K W C K E A K E
 2404 gatcgaaactttccacactcatatacaaatatgggtcaatgggtatgctttggaaaatttgggtgggaagttaattacacacaaca
 85 D R T F S T L I S N M G Q W Y A L E I C W E V N Y T T
 2488 aaatcaggtaaaacgaaaaagagaatctcgaaacaataatttatgatagccttaaaaaataatcctttccagtgaaacaaatt
 113 K S G K T K K E K S R T I I Y D S L K K Y P F P V K Q I
 2572 gcagaagcttttaattttctataaaaaaggcgaaatagattatacaaaaagaaagacatttggttacaacacaaacaaagat
 141 A E A F N F P I K K G E I D Y T K E R P I G Y K P T K D
 2656 gaatgggagttttaagaacgacattcagattatggcgatggcattaaaaattcaattcgatcaaggactaactcgaatgact
 169 E W E Y L K N D I Q I M A M A L K I Q F D Q G L T R M T
 2740 agaggaagcagcgttttagcgattacaagatttggttaaaagctacacatggaaaatcaactttcaacaaatgggtttcctatt
 197 R G S D A L G D Y K D W L K A T H G K S T F K Q W F P I
 2824 ttgtccttttaggggttgataaagacttacgttaaagcatataaaaggcggttccacttgggttaacaaaggttttcaagggaagaa
 225 L S L G F D K D L R K A Y K G G P T W V N K V F Q G K E
 2908 ataggtgacggcattgtctttgatgtcaactcttctcctctcaaatgtacgttaagacctttaccatctggaacacctcta
 253 I G D G I V F D V N S L Y P S Q M Y V R P L P Y G T P L
 2992 ttctacgaaggagaatataacacgaacaacgactatccgctgtacattcaaaatatacaagtaagattccgctttaaaggaggggt
 281 F Y E G E Y K P N N D Y P L Y I Q N I K V R F R L K E G
 3076 tatattccaaccattcaagtttaagcaagttcattattcattcaaaacgaatatcttgaaatcaagtgtaacaagtttaggagtt

319

309 Y I P T I Q V K Q S S L F I Q N E Y L E S S V N K L G V

3160 gacgaattaatcgatcttactcttcaaaatgttgacctagaattatcttttgaacactacgatatttttagagatacattacact
337 D E L I D L T L T N V D L E L F F E H Y D I L E I H Y T
3244 tacggatatatgttcaaagcttctgtgatgttcaaaggctggatcgataaatggatcgaagtaaagaacaccaccgaaggg
365 Y G Y M F K A S C D M F K G W I D K W I E V K N T T E G
3328 gctagaaaaagctaacgcaaaaggtatgttaaatagcttctgtatggaaagtccggaacaaacctgacattacagggaaaagtgcct
393 A R K A N A K G M L N S L Y G K F G T N P D I T G K V P
3412 tacatgggagggacggcattgttcgattgacactaggagaagaagaattaaagagatcctgtttatgttcgcttgcgtatgttt
421 Y M G E D G I V R L T L G E E E L R D P V Y V P L A S F
3496 gtgacggccttgggtagatatactaccattacaaccgctcaaaaatgttttgatcgcatattttatgtgatacagatagcatt
449 V T A W G R Y T T I T T A Q K C F D R I I Y C D T D S I
3580 catctagtaggaacagaagttccagaagcaatcgatcacttggttgatcctaaaaaacttggttattgggggcatgaaagcaca
477 H L V G T E V P E A I D H L V D P K K L G Y W G H E S T
3664 tttcaacgagcaaaattcattcggcagaaaacatacgtagaagaattgatggcgaattaaatgtaaagtgtgctggtatgcc
505 F Q R A K F I R Q K T Y V E E I D G E L N V K C A G M P
3748 gatcgaataaaagagattgtaacttttgacaattttgaagttggtttttcaagctatggaaagtgtgctacctaagaacacaa
533 D R I K E I V T F D N F E V G F S S Y G K L L P K R T Q
3832 ggtggcggtgattagtagacacaatgtttacaatcaataa 3873
561 G G V V L V D T M F T I K *

1820RF003

11305 atggaagaacgaattgatattcaaatgaacaagatgaagaagaaaatcaaaagaattacctattgcacctgaaacgaacccg
1 M E E R I D I Q M N K M K E E N Q K N Y L L H P E T N P
11389 aaacaagttggttttgcattgaaacattgcattgaaatcaaaatcaggagagtttcaacaattttgtgacacaagaaaatgaca
29 K Q V V P D E T L H G N E N Q E S F N N P V D T R K M T
11473 actacaattgatgtaagtgccttatgggttatcgctgacggtgtaacagattgtacaccaatattaaataaattacttgaaag
57 T T I D V S A Y G V I A D G V T D C T P I L N K L L E E
11557 aaaagcgaaatgggtatcacttttttctctcctgtggaacgtgattcatattatcgctttgctaaccattgaaatgaaa
85 K S E M G I T F Y F P P C E R D S Y Y R F A N T I E L K
11641 cgtgatgtacctgtagttactttcttaggatcgggagaaacgacattaaagtgtgaaacaatgacggcatttaattgtaaacatc
113 R D V P V V T F L G S G E T T L K F E T M T A F N V N I
11725 gaaagtttcaatattgatggttttgcatattggttgccacaaaggcgctcaaagtggtaaggaattttcttaatgatattcgc
141 E S F N I D G F A L W L P Q G A Q S G K G I F F N D T R
11809 aattacaatcgcttttgactttgtattgttgcgttaacttttaaatgaaggaacgtatgttggtagtagaggtaga
169 N Y N R F D F D L F V R N C T L N E G T Y V V V A R G R
11893 ggggttacatttgaaaattgtctattctctaatatctctcaagcaattatcaaaacagcttttcccgatgtaaatggtatgtgg
197 G V T F E N C L F S N I S Q A I I K T A F P D V N G M W
11977 caaggggaacgatatacaactagggttacaggttttagaggtttcttgtgaaaaacaacgctattcattttgtacacgcatc
225 Q G N D I N T R G T G F R G F F V K N N R I H F C T A I
12061 attatcgacaatgacgatgattatcagaatgtaatttaattctgtgaaatttctggtaacacaatcgaaaggtgaggttaatt
253 I I D N D D D Y Q N V I N F C E I S G N T I E G G V S Y
12145 tatcgagatagcgataacttgcatgtccaaaacaacacattttctagcatagcgaatagaaacgctttgtttgagttt
281 Y R G Y A H N L H V Q N N H F L A Y G N R N A L F E N
12229 caagatggtgacgaacttatattgatgtagatttctgtgtaactcacaagtcgaggggaatgtagacagattttca
309 Q D V D Q A Y I D V D V Y C R N S Q V E G M N S T A I S
12313 cgtttaattgtttgacggacattaccgaaacttaagattacaggttaattatcgtttgcaaggacatgttatcagttg
337 R L I V V Y G H Y R N L K I T G K L Y R C Q G H V I T L
12397 tatggcggtggcgttaatttctattgtgacttgatggcacaagaacaccttgacggacggttaccggtttattcaaaaggct
365 Y G G G V N P Y C D L M A Q E A P L T D G Y R F I Q T A
12481 gacaatcgagtttaactatgatgggtttgttgcgtggtttgtctaattcaacaaaagtaaatacccaatgattctataagca
393 D N R V N Y D G F V V R G L S N S T K V N T P M I Y K A
12565 cctcagactgttttctataatcgtagaatcgatcatgtgctaacagggtccaaatgcaagtaattgtatataactag 12639
421 P Q T V F Y N R R I D H V L T G P N A S N V Y N *

1820RF004

4626 atggcgtgacaaaatcacagaacaagatgttcttcgtgccacaaatgtagaaacaccagtagaattaatgactgctatttataat
1 M A D K I T E Q D V L R A T N V E T P V Q L M T A I Y N
4710 agttcatcatctcttttccaggcgaacgtacattatgcacaaatgcagataaacatcgaaagcgttggtgacgggatcacagttta
29 S S S L F Q A N V P M P N A D N I E A V G A G I T R L
4794 gacgtagtataaaacgaatttatttcaactttagttgacgctattggtaagtagttatccgatacaaatcttggcgtaaccc
57 D V V K N E F I S T L V D R I G K V V I R Y K S W R N P
4878 ttgaaaatgtttaaaaaaggaaacatgcctttaggtcgaacgattgaagaatttttggtagacattgcacaggaacataagttc
85 L K M F K K G N M P L G R T I E E I F V D I A Q E H K F
4962 aaccctgacgagctgtttacaggggtattttaacaggaagttcccgatgtaaaaacattgttccacgaattaatcgtaaggt
113 N P D E S V T G V P K Q E V P D V K T L F H E I N R E G
5046 tactacaacaaacgatccaagaagcatggttagaaaaagcatttacttcattgggataatttcaatagtttcgtgctggtga
141 Y Y K Q T I Q E A W L E K A F T S W D N F N S F V A G V
5130 atgaacgctttatcacaggtgacgaagtaagcgaatttgaatacacgaaatttataatagcaaaactaccaagaaaagagcta
169 M N A L Y T G D E V S E F E Y T K L L I A N Y Q E K E L
5214 ttcaaagagatcgaatttggcgaattactgaatcaaatgcaaaagaattttatccgtaagatcaaatcaacctctaacaattta
197 P K E I E I G E I T E S N A K E F I R K I K S T S N K L
5298 gaatttatgattccgcttacaacgctcaaggagttaaaacatctacctcaaaatctgatcaatcagttatcagtcgacgac
225 E F M S S A Y N A Q G V K T S T S K S D Q Y V I I D A D

5382 acagacgcaaccattgacggttgacggttttagcagcggcattcaatatgagtaaaactgactttgtaggacacaaaaatcgttatt
253 T D A T I D V D V L A A A F N M S K T D F V G H K I V I
5466 gatgagtttcttaaaaaagaggcgaagaatcgtaaatattgtggcagttattgtatagtagtaaggtttatgatctacgac

320

281 D E F P K K E G E E S S N I V A V I V D S E W F M I Y D
5550 aaattgtacaaaacaagtcctacacccctgaagggttatattggaattattggtgaccaccaccaactatattctact
309 K L Y K T T S L Y N P E G L Y W N Y W L H H Q L Y S T
5634 tctcaattcgggaacgctgttctgtttaaatacagcaaaaacctgtcacaaaagtgtctttgcaagtgcacaactagt
337 S Q F G N A V A F V K S A T K P V T K V A F A S A T T S
5718 gttgttaaaggatcatcctaagatatcgcatgtacatttaccagtagaagcaacaacaacgaaggaagttgtttcatca
365 V V K G S S K D I A L T F T P V E A T N Q Q G E V V S S
5802 gcaccagcattggttaaggcaacgtaaaacaacagcaggttaaagcgaactgccgttaacgtagaaggttagaagtcggtcaa
393 A P A L V K A T V K Q T A G K A T A V T V E G L E V G Q
5886 tcattagtaacattcacagctatcgaggtcaacaagcaacggttcttctgtacggttacttctgactaa 5954
421 S L V T F T A I G G Q Q A T V L V T V T S D *

182ORF005

12651 atggcaactcttacaatgaacaatagcttagaggacaacaatcgctaaaatactttcaaatatggctataataaaaattca
1 M A T L T N E Q I A R G Q T I A K I L S K Y G Y N K N S
12735 caagttaggagttgtcgccaatctccattgggaatcggtgttgaacccgaacagcaatgaatattggtggaggcggtatggg
29 Q V G V V A N L H W E S A G L N P N S N E Y G G G G Y G
12819 ttaggccaatggagcctaaagcaatcttctatcgccaagcaaatcttggtgtgtctaatgctaagctgaacggtggaa
57 L G Q W T P K S N L Y R Q A Q I C G L S N A K A E T L E
12903 ggtcaagcagagatcatcgctcaaggggataaaacaggtcaatggatggataatacacctgtttcttctgcaggttataactaac
85 G Q A E I I A Q G D K T G Q W M D N T P V S S A Y T N
12987 cctcagacccttccagcattttaaacaatctgcaaatattgattgtgtctacaattaattttatgtgtcactgggaacgctcgtt
113 P Q T L S A F K Q S A N I D V A T I N P M C H W E R P G
13071 aaacttcataatcgaagaaagacttgatcttgacaaagcttatagtaagcatattgacggttagcggtggcgtaaaacgt
141 K L H I E E R L D L A Q A Y S K H I D G S G G G G V K R
13155 tgcattggaaacccaatcaagaatacaaatcttgctcctaaaagtctcatgagtggacaacttttggcagcagcaggaac
169 C Y G T P I K N T N L D P K S F M S G Q L F G T H A G N
13239 ggcagacaaataatttccatgatggttggatttgggttcaattgatcaccctggcaatgaaatgattgcatgttgcgtagga
197 G R P N N F H D G L D F G S I D H P G N E M I A C C D G
13323 acagtaacacatggttgaacaatgggagcatttaagagcgtattttgtgataaatgatggtacttacaatatcggttatcaagaa
225 T V T H V G T M G A L R A Y F V I N D G T Y N I V Y Q E
13407 tttagttataaccagtcataataaaggttaaagttggcgacaaagtaagaacggacaagtttgcgcaatcagtgacgggat
253 F S Y N Q S N I K V K V G D K V K N G Q V C A I R D A D
13491 catttacatttaggttttactaaaaaagattttatgactgcgttaggatcttcttcatagatgatggaacatgggaagaccct
281 H L H L G F T K K D F M T A L G S S F I D D G D G G
13575 ttgaagtttttagggcaatgttttgagatggagatactggcgagataatgacgataacaataaggataaaaatgatcttatt
309 L K F L G Q C F G D G D T G G D N D D N N K D K N D L I
13659 tatctattgctatccgatgccttgaatgggtggaatttttaa 13700
337 Y L L L S D A L N G W K F *

182ORF006

14995 atgacaaatagcttaggcgttaaacttgaagagaaaaacttatactataaccctaacaatgcttttaggttttaattgcctaagt
1 M T N S L G V K L E E K N L Y Y N P N N A L G F N C L M
15079 ttgtttgtaaataggcgacgtggtataggtaaaacttatggttataaaaatttgggttaactcgctttattaaacacggcgaa
29 L F V I G A R G I G K T Y G Y K K F V V N R F I K H G E
15163 caattttatttttaagaagattcaaaacagaacttaaaaagattcctcaatttttcaaaacatggcgaaagaatttctgat
57 Q F I Y L R R F K T E L K I P Q F F K T M A K E F P D
15247 cataaacttgaagtaaaaggaaaagaattctattgtgatgataaattaatgggttgggtgttccacttagtagctgggaatt
85 H K L E V K G K E F Y C D D K L M G W A V P L S T W G I
15331 gaaaaatctaataatccccgaagttctgacaaatttggatgagttttaaattgagaaatcaaaaactcattatttacc
113 E K S N E Y P E V R T I L F D E F L I E K S K I T Y L P
15415 aacgaagctgaagccttattgaacatgatggaacgggttttccgaagcgtacaaatacaagatggttattgttgagtaatgca
141 N E A E A L L N M M E T V F R R R T N T R C V M L S N A
15499 actagtgtagtgaaaccttatttctgtatttcaatctgcagccagatttgaataagcgttttaactctatcaagatcgaggt
169 T S V V N P Y F L Y F N L Q P D L N K R F N L Y Q D R G
15583 atattgattgaattgtgtgattcaaaagactttgcagaagtgaaagagaaaacaccttttggtagattgattcggtggaacagaa
197 I L I E L C D S K D F A E V K R E T P F G R L I R G T E
15667 tacgaagatttttagtatcaacaatgagtttgcattgatagtgatacgtttattgaaaagagaagtaaaatagtagtttctta
225 Y E D F S I N N E F V N D S D T F I E K R S K N S S F L
15751 tgcgccattgttttgaagggaaaatctttgggtattggatagacgtgaaacaggttgtgtctatgtgagttgattatcaa
253 C A I A F E G K I F G Y W I D A E T G C V Y V S Y D Y Q
15835 ccaatacaaatcatttttgaatgactacgaaagaccatgaagaaaatagattgctgatgaaaatttggcgaataattat
281 P N T N H F Y A M T T K D H E E N R L L M K N W R N N Y
15919 tatctttcaacagtgccgaagcattcaagaatagtttatctgcgttttgataacattgttattaagaatttaccattatgattg
309 Y L S T V A K A F K N S Y L R F D N I V I K N L H Y D L
16003 ttttaataagatgaaaatctggttaa 16026
337 F N K M K I W *

182ORF007

7795 atgagtagacgaaaagggtgcaggacttgctagaaaataaccgttatcacagcaaaaagcagaccttatccaatgaaccctattca
1 M S R R K G A G L A R N N R Y T A K S R P Y P N E P Y S
7879 agttagtagaagaatcagctactatgaacattatcgtagacaactcacgctccttacggtttcagttgtttgaatgggaaaat
29 S D V E E I S Y Y E H Y R R Q L T L L T F Q L F E W E N
7963 ttgccaataatcaattgacctcgttatttagaaattgctttacacactaatggttatcttgggtttctttaaagacctacactt
57 L P K S I D P R Y L E I A L H T N G Y L G F F K D P T L
8047 ggggttcaggttgcgcaggggagaagatggtcaaatcgatcattatcacacccctattttctttacagcaaacgaagcaatg

321

85 G F M V C A G A E D G Q I D H Y H N P I F F T A N E A M
8131 tatcacaagagatatcctgttttaagatatgatgatgataaatcaaaatgtatcatgttgataataatgacttgaaa
113 Y H K R Y P V L R Y D D D D D K S K C I M L Y N N D L K
8215 gttcctacgttaccaggtttacatcggtttgttttagatatggcgacataaaccagatatcacgagtgaaatcgagagcgcaa
141 V P T L P S L H R F A L D M A D I N Q I S R V N R R A Q
8299 aaaacacctgtaattattcaactgatgaaaagaatacttctcattgctacaagccttataaccaaattgacgaaaataatcag
169 K T P V I I Q T D E K K Y F S L L Q A Y N Q I D E N N Q
8383 gctgtttttgtgataaagatatggagtttgacgaatccttttaattgtatggcaaacaaatgctccatatgtagtagataaacta
197 A V F V D K D M E F D E S F N V W Q T N A P Y V V D K L
8467 cgatcagaattgaacgaagtatggaatgaagtgttaacttttctaggtatcaacaatgctaacgttagataagactgacgtgtg
225 R S E L N E V W N E V L T F L G I N N A N V D K T A R V
8551 caaacatcagaagcttattcaacaatgaacagattgaaagttcaggtaacatcttgtaaaatcaagaaaagagtttgcgat
253 Q T S E V L S N N E Q I E S S G N I L L K S R K E F C D
8635 cgtgtaaatcgtgtctttggcgatgaacttgacggaaagattgacgtgaagtttagaacagacgccttcgacaattacacatg
281 R V N R V F G D E L D G K I D V K F R T D A V R Q L Q L
8719 gcggcaggtcaatcaaaaaagaccagatgagtgagggttgccaagtgtacttaa 8775
309 A A Q S K K D Q M S G G L P S A T *

182ORF008
14105 atgatgaatggattgatattctctagttatcaaacagggaattgatctttcaaaagttccatgcgattttgttaaatattaagca
1 M M N G I D I S S Y Q T G I D L S K V P C D F V N I K A
14189 acaggcggaacaggttatgtaaaccttgattgtgacaggcatttcaacaagctttgtcttttaggttaaaaagattggtgtgat
29 T G T G Y V N P D C D R A F Q Q A L S L G K I G V Y
14273 cattttgcgcagatgagaggggtttagaaggtacacctcaacaagaagcgcaattcttttagataaatattaaggttacattggt
57 H F A H E R G L E G T P Q Q E A Q F F L D N I K G Y I G
14357 aaagctgttcttattcttgactttgaaggtcaaatcagaagatgtaaatggcgaaagcatttcttgattgttttaaat
85 K A V L I L D F E G S N Q K D V N W A K A F L D Y V Y N
14441 aaaacaggcggttaagcattggtttatcgtatcacgaaacctcaatacaactgattttctagtagtcaaaagcgatgt
113 K T G V K A W F Y T Y T A N L N T T D F S S I A G G D Y
14525 gggttatgggttgctgaatatggatcaaatcaaccacaaggctactctcaaccagcgccacctaatacaataattttccaatt
141 G L W V A E Y G S N Q P Q G Y S Q P A P P K T N N F P I
14609 gttgctgttttcaagtttacaagtaaggacgtttaccaggatacaacggcaatcttgatttgaattgtttctatggcgatggt
169 V A C F Q F T S K G R L P G Y N G N L D L N V F Y G D G
14693 aatacatgggatctgtatgtaggttaaaaaacaggatcaaatgttctctctgaaaaataaataatttgacgccacaagtgtgag
197 N T W D L Y V G K K Q D Q I V P P E N K I F D A T S D E
14777 tttatttctactcttacaacaggttagcacagcgtgttttatttgacggagaaacgatctttgaattgtctgacccaacacaa
225 F I P T L T T G S T S V F Y F D G E T I F E L S D P T Q
14861 ctgcgatcatattagaggaacatacaatcatgttcatggaaaagaatcccatcaatgggtgtggacacctgaacaatttgatatt
253 L D H I R G T Y N H V H G K E I P S M V W T P E Q F D I
14945 tacttaaaaatgtatgaaaagaaccagtatataaatag 14983
281 Y L K M Y E K K P V Y K *

182ORF009
8765 gtgctacttaaacgttatattgaaagtttactttattaccaacctgaattatctcgaagaagcgtattgagttggcgcaaaa
1 V L L K R Y I E S P T Y Y Q P E L S R K E R I E V G R K
8849 caattgtttgattttgattatccgttttatgacgaacaaaacgagcagaatttgaacaaaatttatcaatcacttttacttg
29 Q L F D F D Y P P Y D E T K R A E F E T K F I N D N R N K
8933 agagagataggctcagaacagatgggatcatttaagtttaattcttgacgaatatttaaatctaacaatgcctcattggaataaa
57 R E I G S E T M G S F K F N L D E Y L N L N M P Y W N K
9017 attgttcctatcaaatcttgaaagatttccgatttttgatgacatggatcacaccattgatgagaaacagaaattgttaaatag
85 M F L S N L E E F P I F D D M D Y T I D E K Q K L L N E
9101 attgatacaaacatcaaaagcgaatcgtgatgaatcgaagaacaaacgaagcagtagatcaaacagacacagaaacaaaaat
113 I D T N I K A N R D E S K N Q T K Q V D Q T D N R N K N
9185 acacgtgacacaggaacaaacaggttcttctcaaggaaacttatcacagacccccctaaaaagatttgagaattgccaagcaat
141 R D T G T T D S F S R N T Y T D T P Q K D L R I A S N
9269 ggagatggaacaggtgtaataatcattatgcaacaaatatcacagaagatttgagtaagaaacaaagctccacagggcgttgaa
169 G D G T G V I N Y A T N I T E D L S K E T T S T S T G V E
9353 acaaaacacgacaaaacaaatcaaaatacacgaagcaatgcttctgaaaaagaaacaaagaacacagacattaataaagatcaa
197 T N N D K T N Q N T R S N A S E K E T K N T D I N K D Q
9437 aatcaaaccaagatagcattacagatataaaggtaaaagggaaacactgattatgctgacttactcgaaaaatattcgtaga
225 N Q T K D T I T R Y K K G N T D Y A D L L E K Y R R
9521 agtgttttgagaattgagaaaatgatcttttagagaaatgaacaaggaaggcttatttctccttgtttatggagggaggtag
9601
253 S V L R I E K M I F R E M N K E G L F L L V Y G G R *

182ORF010
1310 ttgaccgtaagaatatcaagaatgatagagccaagtttagagaaaatctacggtaaatctaacaagctcgtaaaaaatacaat
1 L T V R I S K N D R A K L E K I Y G K S N K A R K K Y N
1394 cgtttaagacaaaaaggagttgaggaaggcaacttccaactgttccaacatcaagaaaagactttgactacgtataaaatca
29 R L R Q K G V E E R Q L P T V P T S K K R L I D Y V K S
1478 acaaatatgagtcgtagtattttaacaagatgttagacgagttggttagattttgcacaaccttacaacgagaaatttaatttt
57 T N M S R S D F N K M L D E L V D F A Q P Y I F
1562 gagatcaaacagcgaattgttgaattctcaagcgcaaatcaagaagcgcaaattaaaacagagcaagctcaaaaaagcgaaa
85 E I N K R N V A I S R A Q I K E A Q I K T E Q A Q K A K
1646 gaagaacactacaagagcttaacaaagttgaagtttaagaagccacagaaaacaaatgttcacaccaactattttaacagag
113 E E H Y K E L N K V E V K K P T E N T I V T P T I L T E
1730 ttaggtgctgacttaccttttcaagcaatcacagatttttaattatgacgctttcacttctccagaaggagttcagttctattta

322

141 L G A D L P F Q A I P D F N I D A F T S P E G V Q S Y L
1814 gaaaatataggaacaagacgaacaatattttgacgaaagagaccaactttattacgacaatttcagacaagcgatgtttact
169 E N I G K Q D E Q Y F D E R D Q L Y Y D N F R Q A M F T
1898 attttcaattcagacgctgacgatattgttcgtttacttgactcaatggggttgatctatttatgaaacatatgttagtaac
197 I F N S D A D D I V R L L D S M G L D L P M K T Y V S N
1982 ttcttagacatgaaccttgactacattttgacgaagcagaagtacaacagaaaaagaacaagtttacagtaagattgcaaaa
225 F L D M N L D Y I Y D E A E V Q Q K K E Q V Y S K I A K
2066 gtgacgagctgaaacaggtggagaagtcctccctcatataacccacgaagaacatcacataattcagaacacgaggaagaa
253 V I E S E T G G E V P S Y N P T K N I T I N S E T G E E
2150 ttatga 2155
281 L *

182ORF011
9607 atggtagattttaaccccgacaagcggtttgacgggtttaccgctgtattcaaagaacgcttttagcaaatatcctcactactgaa
1 M V D P N P D K R F D G L P A V F K E R F S K Y P H T E
9691 tacagatatgaattactattagatgaagaagtatcggtttaattgcctatctgaatgaagttgggtcttagttaatgatg
29 Y R Y E L L L D E E V S A L I A Y L N E V G A L V N D M
9775 agtggttatttaattactttatcgaacattttgtgagaagttagaagagatcacaaatgacacactcaaaaaatgggtgtct
57 S G Y L N Y F I E H F V E K L E E I T N D T L K K W L S
9859 gatggtagcttagaaaaatcaatgatgactgttttgcgaattatcaaagaatcaaaagattacaaattcttggtgtct
85 D G T L E N L I N D T V F A N Y I K E I K R L Q I L V A
9943 gaaacacgtgctaacagtgatgaattcttttgacaaaaataaacggatgttgctgatgacgaacattttggtataagatt
113 E T R A N S V N I L T K N K P D V A D D R T F W Y K I
10027 caacgcgacaactactgattatggagccgatcctattgacacgttacgtattgttgcaatcaataaagttagtggctggaatacc
141 Q R D N T D Y G A D P I D T L R I V A I N K V S G W N T
10111 gctacaggagatatttatcttaacattaaaggaacggggtgtataa 10158
169 A T G D I Y L N I K G T E G V *

182ORF012
10872 atggcaataaaaaatattcaaatgaaggatagcaatgacaataatttatccaagtgttcgagcagaaaaactgttagatttg
1 M A N K N I Q M K D S N D N N L Y P S V R A E N L L D L
10956 accagtcgtgctgaattacaatgacaaattgtcaatttatatgcagctggtgataaaacaaatgcaattcttctatcgggtgca
29 T S R A E L T M T N C Q L Y A A G D K T N A I
11040 gtaggtatgctcgaaggtatgataaagtttactgaaagtttgacaaacccgtgatcacacgctaccagaaggttttagacca
57 G M L E G M I K F T E S L T N P V I T T L P E G F R P
11124 ataagaacaaaacgtattggtgttctgcgaataattacacacaaatccaacagatacaaaaagaatgggttatgtatcaatc
85 I R T K R I G C F A K Y Y T P N P T D T K E M V Y V S I
11208 acacgtgatggcaagtaactgtaaatgacaatgtaggtaaaatcgaatatctatccctagataaattgcgttttccctctaaaa
113 T P D G K V T V N D N V G K I E Y L S L D N C V F P L K
11292 taa 11294
141 *

182ORF013
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1 M A D K N I Q M Q D K D H N R L M P V T I A K N V L T G
10540 gactctaattctgaattagttaatgctgaaataagaggttaacgctagtgaagctaaaacacttgacacaacagctaaagaaact
29 D S N L E L V N A E I R G N A S E A K T L A Q Q A K E T
10624 gctgctggtttgtcaacagaaattgacacagtaaacacgcaaatcaagcgttgacgaaggtggtacagcacaacaaacc
57 A A G L S T E I D T V T S T A N Q A L T K A G T A Q Q T
10708 gcagaacaagcgaacaaacagcaaacagtatcagcgcagttgcaacggcagctaaaaacacagctgactcagcacaacaaagtt
85 A E Q A K T T A N S I S A V A T A A K N T A D S A Q K S
10792 gcaactgatctagctgttcgagtaagcagtttagagggacacagcaatacataatactgtattaccatag 10860
113 A T D L A V R V S S L E D T A I Q Y T V L P *

182ORF014
13716 atgatagaatatcacacaatggttggcagatgataatcatcttgtttatggtttgatttatatggtttaatggttgcaatgatt
1 M I E Y I T Q W L A D D N H L V Y G L I I W L M V A M I
13800 atcgattttgtgttaggttttacaattgccaaatttaacaagggaatcgacttttagtagttttaaagctaaaagcaggtatcatt
29 I D F V L G F T I A K F N K E I D F S S F K A K A G I I
13884 gttaaggtggcagaatgggttttagtgggttactttatctcgtgtagcagtaaaatcggtgtaggttagttacaaatgtatata
57 V K V A E M V L V V Y F I P V A V K F G A V G I T M Y I
13968 acaatgttgggttgggtttgattttatcagaatttatagatactaggacatatttcagatatcgatgatgataaataattggact
85 T M L V G L I L S E I Y S I L G H I S D I D D N N W T
14052 gattatgttaagaagtttttagacggaacactcaacagaaaggacgatattaaatga 14108
113 D Y V K K F L D G T L N R K D D I K *

182ORF015
854 atggaaatcgtaaaaagcacatttgacacacaaacaccagaaggaatgttacaagtattcaatgccacaaacgggggttcaatt
1 M E I V K S T F D T Q T P E G M L Q V F N A T N G A S I
938 ccgttacgtaacgcaattggcgaagtactagaattgaagatattctagtttactcagacgaagtttctggttttggtggagcc
29 P L R N A I G E V L E L K D I L V Y S D E V S G F G G A
1022 gaaccatcacagcagaactagtcgctttcttcacagaagatggtaaaactttatgcgggtgtatcagcagtagcaacaaatca
57 E P S Q A E L V A F P T E D G K T Y A G V S A V A T K S
1106 gctaaaaacctaattgatgatgactgctaaccctgacatcaaaacaaaaatttctttgtcgaaggaatcaaacgggtgga
85 A K N L I D M M T A N P D I K P K I S F V E G - K S - G
1190 caaaaatttgaataatctacaagtgggttctactgtag 1225
113 Q K F V N L Q V V S L *

182ORF016
17033 atgattaacaatttatcattaatttttagaggggttaaaactaaactaaagatgacaacgatagtttagcgtctatcaagtca
1 M I N N L S L I L E G L N Q L T K D D N D S L A S I K S
16949 gaaataacacaaggaggaaaacattaattttatattgattacgtttacaaaagagttcgtgttaacacatgataataaac

323

29 E I T Q G G K Q L I L Y I D Y V T K E F V L T H D K Y N
16865 tatgtttatcttgatagccattgcattaatcgcaataacgaaatcaatgaaaagcgttgaaactatgcggaacaattgaaa
57 Y V Y L D S H C I N I A I T K S M K S V E H Y A E Q L K
16781 catgacggatataaacaattacggacaatag 16749
85 H D G Y K Q I T D K *
182ORF017
154 atgaaatattcactacaacaatagatgaaatcaaatcaaatcttcagaattagattaaaaagcgtgaactagaggaattg
1 M K Y S L Q Q I D E I K S T I F R I R L K R H E L E E L
238 gtggacgaagtaaacgatattgctaagatccggaggaaagatatcttttatcgttttattacagagaagaacgtttgttt
29 V D E V N D I A K D P E E R Y L L S F Y Y T E E E R L F
322 gaaattccctctgcaagattaatagattattacaagaaaagatcacaatctgaaatcggaatcatatcactcgaaaaaaga
57 E I P S A R L I D Y Y N E K I T N L K S E I I S L E K R
406 ttacaaaaactagtaaaataa 426
85 L Q K L V K *
182ORF018
16737 atgattgcacgaacattcaagaacaccgcgaactaattgaatggttacgtttctactgtaaacgtaacctttcagacaatgaa
1 M I A R T P K E H R E L I E W L R F Y C K R N L S D N E
16653 aaaatagagatcatagaggggactttacaagattcgacgttccggaataaatatcacggaactttgttaactcattcaacg
29 K I E I I E G T L Q D F D V P E I N I T E L L L T H S T
16569 ctattaccggaatcgagtcatttaacattcttgaaaggtattgtcaggcaatgaaattagtaacttcatacgtaaaaagttggt
57 L L P E S S Q F N I L E K Y C Q A M K L V T S Y V K V G
16485 tctcgctatcagttagcgttacaaaataccaaaaggctatttaaggaggtggaataa 16429
85 S R Y Q L A L Q I P K G Y L K E V E *
182ORF019
4323 atgaaaattaaagaacatgaatcaatttttaattggtattcttgaaagtgtcacagacgggtgaagcaagatcaagattgtagaa
1 M E I K E H E S I L N G I L E S V T D G E A R S K I V E
4407 catcttgaagcattgcgagaagactacggagcaacaactgaagctttgacatcagcaaatagcacacttgaaggttaaaagaaa
29 H L E A L R E D Y G A T T E A L T S A N S T L E K L K K
4491 gataacgaagcgttggtttttcaactcaaatgttccgagaacgagcgatcgtagaaccagcagaaaaataacgaaccagaa
57 D N E A L V I S N S K L F R E R A I V E P A E N N E P E
4575 acagaccagaatattacactagacgatttaggaatttaa 4613
85 T D Q N I T L D D L G I *
182ORF020
10158 atggcagacattagaacacaactaacaagtgaagatggatcagacaattttttccaatttcaaaagccgttaattattatgact
1 M A D I R T Q L T S E D G S D N L F P I S K A V N I M T
10242 aatagcgggtacgaatgtagaaggagaattgggtacactcaaaacaaatgacgaacaaatgaatacctcagttcaaatgctgta
29 N S G T N V E G E L G T L K Q N D E T M N T S V Q N A V
10326 gttactgccaatcaagcaaaagattctgtagctgaatataatgtaaatgttggttaactaaccatcgaaataacaacattagag
57 V T A N Q A K D S V A E L N V N V G K L T N R I T T L E
10410 agtcagtggtgaatcttggttatctgtatgtagaggtgtaa 10454
85 S T V A N L D G I R Y V E V *
182ORF021
17339 atgaacaataaatcattaatagctgaaaaaggagaggtatctctacttcacccctttaatgagtggtgatgaattatcatatc
1 M N N K S L I A E K G E V S L L H P F N E W D M N Y H I
17255 atagataccgaaaacaataaacattatcttattgataatgtaggtaggcgatgaggaatattgtttgttatcttttgaagaa
29 I D T E N N K H Y L I D I D E V G D E E Y C L L S F E E
17171 ctaaaaggaattagataggtatcttattccgagttatcatggaataacacagaaataacatattaa 17106
57 L K E L D M D L I S E Y S W K T T E I T Y *
182ORF022
12868 gtgggtgtgtctaatgctaagctgaaacgttggaaggtcaagcagagatcatcgctcaaggggataaaacaggtcaatggatgg
1 V G C L M L K L K R W K V K Q R S S L K G I K Q V N G W
12952 ataatacacctgtttcttctgaggttataactaacctcagacccttcagcatttaacaatctgcaaatattgatgttgta
29 I I H L F L L Q V I L T L R P F Q H L N N L Q I L M L L
13036 caattaattttatgtgtcactgggaacgccctggttaaacttcataatcgaagaagacttgatcttgcaagcttatagtaagc
57 Q L I L C V T G N A L V N F I S K K D L I L H K L I V S
13120 atattgacggtagcgggtggcggtgta 13149
85 I L T V A V A V A *
182ORF023
12189 atggtgtgtgttttgacatgcaagttatgcgcataatcctcgataataaacttacgccaccttcgattgtgttaccagaaatctc
1 M V V V L D M Q V M R I S S I I T Y A T F D C V T R N F
12105 acagaaaatttaattacattctgataatcatcgctcattgtcgataatgatcgctgtacaaaaatgaatacgggtgtttttcaciaa
29 T E I N Y I L I I I V I V D N D R C T K M N T V V F H K
12021 gaaacctctaaaacctgtaccttagtattgatatcggttcccttgccacataaccatttacatcgggaaaagctgttttgataat
57 E T S K T C T P S I D I V P L P H T I Y I G K S C F D N
11937 tgcttgagagatattagagaatag 11914
85 C L R D I R E *
182ORF024
6174 atgcttgtaactatctcatctttaaacaagaacttatcctagtaaatggcagtagcctttgttactgatattgaataa
1 M L V T I S S L K T K K L I L V N G S M P L L L I L N I
6258 agaattgacaacaaagttctgttacctttgaattgattgttttacaacttatcggttcgatattggtatagcagaaagtttca
29 R M T T Q V S L P L K L M F Y K L I V S I L V Y E K V S
6342 ttgcaaaagaacacctcaactttattattcgaatggaatacctttcattaatacaattgaagagtcggttgattacggtagag
57 L Q K N T L N F I I R M E Y L S L I Q L K S R L I T V E
6426 aatacacaacaacaatgtaa 6446
85 N T Q Q Q M *

182ORF025

548 atggggtcgaaaactaatgcaacgaaacgtaacatcaactaaagtagaattctcagaagttatcgtaacagatggagcgccaaca
1 M G R K L M Q R N V T S T K V E F S E V I V Q D G A P T
632 attgtaccatgcgaaccagttgtcttaacaggaaaactttcagaagaaaaagctttatcagcgatcaaacgtaaaacccgtgat
29 I V P C E P V V L T G K L S E E K A L S A I K R K N P D
716 aaaaacgtagttgtaacaaatggttcacatgaaacagcgctttacacaatgccagtcgataaaatttcagagtttagcagacaaa
57 K N V V V T N V S H E T A L Y T M P V D K F I E L A D K
800 tcaacacaagcctaa 814
85 S T Q A *

182ORF026

13259 atggaaattatttggctgctgctgttcctgcagtcggtgcaaaaagttgtccactcatgaaacttttaggatcaagatttgtatt
1 M E I I W S A V S C M R A K K L S T H E T F R I K I C I
13175 cttgattgggtttccatagcaacggttttacgccaccgcccagctacgctcaatattgtactataagcttggcaagatcaag
29 L D W G S I A T F Y A T A T A T V N M L T I S L C K I K
13091 tctttctcgatatgaagttaccagggcggttccagtgacacataaaatttaattgttagcaacatcaattttagcagattgtt
57 S F F D M K F T R A F P V T H K I N C S N I N I C R L F
13007 aaatgctga 12999
85 K C *

182ORF027

14896 atgaacatgattgtatgttctctaatatgatcgagttgtgttgatcagacaattcaagatcggtttctccgtcaaaataaaa
1 M N M I V C S S N M I E L C W I R Q F K D R F S V K I K
14812 cagcgttgtgctacctgttgaagtgtaaaataaaactcatcacttggcgctcaaatattttttagcaggagaaacatttg
29 H A C A T C C K S E N K L I T C G V K Y F I F R R N N L
14728 atcctgtttttacctacatacagatccatgtattaccatcgccatagaaaacattcaaatcaagattgctgttattcctgg
57 I L F F T Y I Q I P C I T I A I E N I Q I K I A V V S W
14644 taa 14642
85 *

182ORF028

14430 atgtttataataaaacaggcggttaagcatgggtttatcgtatatacagcaaacctcaatacaactgatttttctagtattgcaa
1 M F I I K Q A L K H G F I R I Q Q T S I Q L I F L V L Q
14514 aaggcgattatgtttatgggtgtgtaatatggatcaaatcaaccacaaggctactctcaaccagcgccacctaataaaata
29 K A I M V Y G L L N M D Q I N H K A T L N Q R H L K Q I
14598 attttcaattgtgctgttttcagtttacaagtaaggagcgtttaccaggatacaacggcaatcttgatttga 14672
57 I F Q L L P V F S L Q V K D V Y Q D T T A I L I *

182ORF029

17606 atgaatgaaccgatcgatatacacagaatttattcaataacgctggtatgtatgaaaatttttagagatgaggataaacttagt
1 M N E P I V Y T E I Y S N N V V C M K I F R D E D K L S
17522 aaattcctctatttagaatttgagtggtgagtgaggctaaaagttacttgaaaataaaacatttcttgatgataactggact
29 K F L Y L E F E V D E A K K L L E N K T I S F D D N W T
17438 ttctcaataaattatccagaatataa 17412
57 F S I N Y P E Y *

182ORF030

16429 atggctacattctacaaggaaccaatatacagatcacagatattttatatagatgggtgggaggttttgatacacaaaaccgaa
1 M A T F Y K E P I Y D I T V F Y I D G W E V L I H K T S
16345 cctctcacttaacaaaagcattaaaatatagccgtatatacctagaaatggatagtgattgaggttagaataagaaagaat
29 P L T L T K A L K Y S R I Y L E M D I V N C V R I E R N
16261 ggacgtcctatagctacattttacaggggaattataaaactgtataaggagaagaactatga 16199
57 G R P I A T F Y R E L L K L Y K E K E L *

182ORF031

8603 atgttacctgaactttcaatctgttctattgttagataagacttctgatgtttgtacacgtgcagtccttatctacgttagcattg
1 M L P E L S I C S L L D K T S D V C T R A V L S T L A L
8519 ttgatacctagaaaagtttaacacttcttccataacttcgttcaattctgatcgtagtttatctactacatattggagcatttgg
29 L I P R K V N T S F H T S F N S D R S L S T T Y G A F V
8435 tgccatacattaaaagattcgtaaaactccatatctttatccacaaaaacagcctga 8379
57 C H T L K D S S N S I S L S T K T A *

182ORF032

11413 atgtttcatcaaaaacacttgtttcgggttcgtttcaggggtgcaataggtaattcttttgattttcttctttcatcttgttca
1 M F H Q K Q L V S G S F Q G A I G N S F D F L L S S C S
11329 ttgaaatcaattctgttcttccatatagaacctcttatttttagagggaaaacgcaattatctaggatagatattcgatttta
29 F E Y Q F V L P Y E P P Y F R G K T Q L S R D R Y S I L
11245 cctacattgtcatttacagttactttgccatcaggtgtgattgatacataa 11195
57 P T L S F T V T L P S G V I D T *

182ORF033

4942 atgtcaacaaaaatttcttcaatcggttcgacctaaggcatgtttcttttttaaacattttcaaagggttacgccaagatttg
1 M S T K I S S I V R P K G M F P F L N I F K G L R Q D L
4858 tatcggaataactctttaccaatacgggtcaactaagttgaaataaattcggtttttactacgtctcaaacgtgtgatccctgca
29 Y R I T T L P I R S T K V E I N S F F T T S K R V I P A
4774 ccaaccgctcgatgttatctgcatttggcatagtcagcttcgcctga 4727
57 P T A S M L S A F G I G T F A *

182ORF034

6160 gtgtttatctactctaaaaactccccgagttgtgtatcccttttgataagaacaatctctattctcgtaagaacaggaaacga
1 V F I Y S K N S P E L C I P L I R T I S I L V K N R K R
6076 attaaagtacgattctctgttctgttgaagtttaaacactcttgggtgtgtataggtgttatcaaaaggcagcttagccaacaa
29 I K V R F L F L L S F K P S C V C I G V I K R H V S Q Q
5992 ttttacatttgataccttcttgcataattgtctctcttag 5951

325

57 F Y I C I P S C H N C P P *
182ORF035
15758 atggcgcatagaactactatttttacttctcttttcaataaacgtatcactatcattgacaaactcattgttgatactaaaa
1 M A H K K L L F L L L F S I N V S L S L T N S L L I L K
15674 tcttcgtattctgttccacgaatcaatctacaaaagggtgttctctcttcacttctgcaaagctctttgaatcacacaattca
29 S S Y S V P R I N L P K G V S L F T S A K S F E S H N S
15590 atcaatatacctcgatcttga 15570
57 I N I P R S *
182ORF036
2315 atgtctgtgctgcttgcattttacaccactcaaaaaagaatcgatttctaaaccgaacgtcatattgtcaacgtgtgtctata
1 M S V L P C I L H H S K K E S I S K P N V I L S T L S I
2231 tcgcatacgccccacgaccatacacgacaatcggttgagatcagttgttcttcaaagtcgccagtattttcttaatacataatt
29 S H T P H D H T R Q S L R S V V V S K S P V Y F L I I I
2147 cttctcctgtttctgaattaa 2127
57 L L L F L N *
182ORF037
12280 gtgagttacgacaataaacatctacatcaatataagcttgatccacatcttgaaactcaaaacaaagcgtttctatttccgtatg
1 V S Y D N K H L H Q Y K L D P H L E T Q T K R F Y F R M
12196 ctgaaaaatggtgtgtgttttgacatgcaagttatgcgcatatcctcgataataacttacgccaccttcgattgtgttaccag
29 L E N G C C F G H A S Y A H I L D N N L R H L R L C Y Q
12112 aaatttcacagaaattaa 12095
57 K F H R N *
182ORF038
14769 gtgagtgagtttattttcactcttacaacaggtagcacaaagcgtgttttattttgacggagaaacgatctttgaattgtctgatc
1 V M S L F S L L Q Q V A Q A C F I L T E K R S L N C L I
14853 caacacaactcgatcatattagaggaacatacaatcatgttcatggaagaaatcccatcaatggtgtggacacctgaacaat
29 Q H N S I I L E E H T I M F M E K K S H Q W C G H L N N
14937 ttgatatttacttaa 14951
57 L I F T *
182ORF039
9992 atgttgctgatgatcgcaacattttggtataagattcaacgcgacaatactgattatggagccgatcctattgacacgttacgta
1 M L L M I E H F G I R F N A T I L I M E F I L L T R Y V
10076 ttgttgcaatcaataaagtttagtggtggaataccgctacaggagatatttatcttaacattaaagggaagggtgtataat
29 L L Q S I K L V A G I P L Q E I F I L T L K E R R V Y N
10160 ggcagacattag 10171
57 G R H *
182ORF040
16202 atgagaaaagatttcgtctacattaacacacccgatccaaaagcaaaacaaaaggcgttagcaaaaatcactaacgccaaagaa
1 M R K D F V Y I N T P D P K A N K K A L A K I T N A K E
16118 ccaaaaacaaactatcgagactacaattactatgttatctactattcatcattgtaatagaactaatcggtgtagctctacta
29 P K Q N Y R R L Q L L C Y L L F I I V I E L I V V A L L
16034 aaatag 16029
57 K *
182ORF041
3886 atggaactatataaagcaatgtttatcgtagctgatgaaggtactattgacggttacgatactgaacactatgtagatatttct
1 M E L Y K A M F I V R D E G T I D G Y D T E H Y V D I S
3970 ttacatgactttgaagaatatatggaagaaacacgtgaaattgaagcagtaacattagtaaaaacaggaaatttaaaaaa
29 L H D F E E I Y G K E T R E I E A V T L V K T G N L K K
4054 taa 4056
57 *
182ORF042
10832 gtgtcctctaaaactgcttactcgaacagctagatcagttgcacttttttgtgctgaatcagctgtgttttttagctgccgttgca
1 V S S K L L T R T A R S V A L F C A E S A V F L A A V A
10748 actgcgctgatactgtttgctgtgtttttcgctgtgttctgcggtttgtgtgctgtaccagccttcgtcaacgcttga 10671
29 T A L I L F A V V F A C S A V C C A V A F V N A *
182ORF043
10652 gtgtcaatttctgttgacaaaccagcagcttttcttagctgtgttgcaagtggttttagcttactagcgttacctcttatt
1 V S I S V D K P A A V S L A C C A S V L A S L A L P L I
10568 tcagcattaaactaattcaagattagagtcgcctgttagaacatttttagcaattgtaacaggcattaaacgattatga 10491
29 S A L T N S R L E S P V R T F L A I V T G I K R L *
182ORF044
6457 atgaaaagttgttacatttgtgtgtgtattctctaccgtaataacgcgactcttcaattgtattatgaagggtattccatt
1 M K S C Y I C C C V F S T V I K R L F N C I N E R Y S I
6373 cgaataataaagttgaggtgttcttttgcattgaaactttctcgataccaataatcgaaacgataagttgttaa 6299
29 R I I K L R V F F C N E T F S Y T N I E T I S L *
182ORF045
6729 atgaatgggtatacctgtatcagcgttatatacatcccgactatcttatttttaaaaagggttcttctgttgtaagaacgccatg
1 M N G I P V Y D V T Y I P T I L F K K G S F V V R N A M
6645 tacttccaaaatttagcattgctgccccattgtgtttgtatatacctccccacttgaattgataggaagtaaaataa 6571
29 Y S P K L A L P A P F G L Y T S P L E L I G S K *
182ORF046
2372 atgggtttcaaatggtgtaagaagcaaaagaagatcgaaactttccacactcatatcaaatatgggtcaatgggtatgctttgg
1 M V S N G V K K Q K K I E H S P H S Y Q I W V N G M L W
2456 aaatttggtgggaagttaattacacaacaaacaggttaaaacgaaaaagagaatctcgaaacaataa 2527
29 K F V G K L I T Q Q Q N Q V K R K R N L E Q *

182ORF047
13353 atgctcccattgttccaacatgtgttactgttccatcgcaacatgcaatcatttcattgccagggtgatcaattgaaccaaagt
1 M L P L F Q H V L L F H R N M Q S F H C Q G D Q L N Q S
13269 ccaaacatcatggaaattatttggctctgccgtttctctgcgtgcgtgccaaaaagtgttccactcatga 13201
29 P N H H G N Y L V C R F L H A C Q K V V H S *
182ORF048
3395 atgtcagggtttgttccgaactttccatacaagctatttaacatacctttggcgttagcttttctagcccccttcggtggtgttc
1 M S G F V P N F P Y K L F N I P L A L A F L A P S V V F
3311 tttaactcgatccatttatcgatccagcctttgaacatatcacaagaagctttgaacatatatccgtaa 3243
29 F T S I H L S I Q P L N I S Q E A L N I Y P *
182ORF049
1578 atgttgcaatctcaagagcgcaaatcaaagaagcgcaaatataaacagagcaagctcaaaaagcgaagaagaacactacaaag
1 M L Q S Q E R K S K K R K L K Q S K L K K R K K N T T K
1662 agcttaacaaagttgaagttaagaagcccacagaaaacacaattgtcacaccaactattttaa 1724
29 S L T K L K L R S P Q K T Q L S H Q L P *
182ORF050
8012 atgggttatcttggtttctttaaagaccctacacttgggttcatggtttgcgcaggggcagaagatgggtcaaatcgatcattatc
1 M V I L V S L K T L H L G S W F A Q G Q K M V K S I I I
8096 acaaccttattttcttacagcaaacgaagcaatgtatcacaagagatatctgttttaa 8155
29 T T L F S L Q Q T K Q C I T R D I L F *
182ORF051
9390 atgcttctgaaaaagaacaaagaacacagacattaataaagatcaaaatcaaaccaagatacgtattacacgatataaaggta
1 M L L K K K Q R T Q T L I K I K I K P K I R L H D I K V
9474 aaaaagggaacactgattatgctgacttactcgaaaatatcgtagaagtggtttga 9530
29 K R E T L I M L T Y S K N I V E V F *
182ORF052
4096 gtgatagttgacaagagtcaaatttggcgagattggcgcaatgtacacgtgaaatcgtgcgctcccgttaagttatggacac
1 V I V D K S Q I W R D W A N V H V K Y R A L P L S Y G H
4180 ataaacgttttgaccgtcaacaaatcgcaaaaaccttttaggagtagcccttaa 4233
29 I N V L T V N Q S Q K P F R S S P *
182ORF053
15656 gtggaacagaatacgaagatttttagtatcaacaatgagtttgcattgatagtgatacgtttattgaaaagagaagtaaaaaata
1 V E Q N T K I L V S T M S L S M I V I R L L K R E V K I
15740 gtatgttcttatgcgcattgttttgaagggaacattcttgggtattggatag 15793
29 V V S Y A P L L L K G K S L G I G *
182ORF054
8136 gtgatacattgtctgttctgtgtaaaagaaatagggtgtgataatgatcgatttgaccatcttctgcccctgcgcaaacat
1 V I H C P V C C K E N R V V I M I D L T I F C P C A N H
8052 gaaccaagtgttaggttctttaaagaaacaaagataaccattagtggtga 8002
29 E P K C R V F K E T K I T I S V *
182ORF055
8324 atgaaagaaatacttctcattgtctacaagcttataacaaattgacgaaaataatcaggctgtttttgtggataaagatatgg
1 M K R N T S H C Y K L I T A K L T K I I R L F L W I K I W
8408 agtttgacgaatctttaaagtgtatggcaacaaatgctccatagtag 8455
29 S L T N L L M Y G K Q M L H M *
182ORF056
6549 gtggcccatctccttttcttatttacttctcatcaattcaagtgagggtatatacaaaccaaatggggcaggcaatgcta
1 V A H L L F P I I Y F L S I Q V G R Y T N Q M G Q A M L
6633 attttggagagtacatggcgtttcttacaacgaaagaaccttttttaa 6680
29 I L E S T W R F L Q R K N L F *
182ORF057
8264 atgtcgcgcatactaaagcaaacgatgtaaaacttggttaacgttaggaactttcaagtcattattatacaacatgatatttt
1 M S A I S K A K R C K L G N V G T F K S L L Y N M I H F
8180 gatttatcatcatcatcatctttaaagcagatattcttctgtga 8133
29 D L S S S S S Y L K T G Y L L *
182ORF058
5176 gtgtattcaaatcgcttacttctgcacctgtgtataaagcgttcattacaccagcaacgaaactattgaaattatcccatgaa
1 V Y S N S L T S S P V Y K A F I T P A T K L L K L S H E
5092 gtaaatgctttttctaacatgcttcttggatcgtttgtttag 5048
29 V N A F S N H A S W I V C L *
182ORF059
15876 atggcttttctgtagtcattgtcataaaaaatgatttgtatttgggtgataatcataactcacatagacacaacctgttccagcgtc
1 M F T R S H C I K M I C I W L I I I T H I D T T C F S V
15792 tatccaatacccaagattttcccttcaaaagcaatggcgcataa 15748
29 Y P I P K D F P F K S N G A *
182ORF060
15404 gtgatttttgatttctcaattaaaaactcatcaacaaaattgtacgaacttcgggatattcattagatttttcaattccccac
1 V I F D F S I K N S S N K I V R T S G Y S L D F S I P H
15320 gtactaagtggaaacagcccaacccattaatttatcatcacaatag 15276
29 V L S G T A Q P I N L S S Q *
182ORF061
2102 atgaggggacttctccactgtttcagactcgatcacttttgcaatcttactgtaaacttgttcttttctgtgtacttctg

327

1 M R G L L H L F Q T R S L L Q S Y C K L V L F S V V L L
2018 cttcgtcataaatgtagtagtcaagggttcattgtcctaagaagttactaa 1974
29 L R H K C S Q G S C L R S Y *
182ORF062
1992 atgtcctaagaagttactaacatatgttttcataaatagatcaagccccattgagtcagtaaacgaacaatatcgctcagcgtct
1 M S K K L L T Y V F I N R S S P I E S S K R T I S S A S
1908 gaattgaaaatagtaaacatcgcttgcctgaaattgtcgttaa 1867
29 E L K I V N I A C L K L S *
182ORF063
14306 gtgtaccttctaaacccctctcatcgcgcaaaatgatacacaccaatctttttacctaagacaaagcttgttgaaatgctcggt
1 V Y L L N P S H A Q N D T H Q S F Y L K T K L V E M L G
14222 cacaatcagggtttacataacctgttcgcctgttgccttaa 14181
29 H N Q G L H N L F R L L L *
182ORF064
7356 atgatgttagtcaaaccaacaaaagggtgttacttgcctaaggctgaaagatcgctcctcctgtactcattgcactgtttccc
1 M M L V K P T K G L L L A K A E K I A P P V L I A L F P
7272 ataccatgtctgaaagtattgcgaatgttttgccttga 7234
29 I P C L K V L R M F C S *
182ORF065
3582 atgaatgctatctgtatcacataaataatgcgatcaaacatttttgagcggttgaatggtagtatctaccccaagccgt
1 M N A I C I T I N N A I K T F L S G C N G S I S T P S R
3498 caaaaactagcaagcggaacataaacaggatctcttaa 3460
29 H K T S K R N I N R I S *
182ORF066
4234 atgtggctactctttttgtgtttcacagaattatgtttcacgtgaaacagtttttatggtataatagaatcaaaaggaggtgg
1 M W L L F F V F H R I M F H V K Q F L W Y N R I K R R W
4318 agattatggaaattaagaacatgaatcaatttttaa 4353
29 R L W K L K N M N Q F *
182ORF067
13882 atgatacctgcttttagcttttaaaactactaaagtcgatttcttgcgttaaatttgcaattgtaaaacctaacacaaaatcgata
1 M I P A L A L K L L K S I S L L N L A I V K P N T K S I
13798 atcattgcaaccattaaccatataatcaaacataa 13763
29 I I A T I N H I I K P *
182ORF068
7267 atgtctgaaagtattgcgaatgttttgccttgcagcaatcaaggagttttgtttccttgcataatgcagaagcatagtcaga
1 M S E S I A N V L L L S N Q G V F V S L H E C R S I V R
7183 ttttaactcctacatcggttaggatcattatcgattaa 7148
29 F N S Y I V R I I I D *
182ORF069
5027 gtggaacaatgtttttacatcgaggaaacttctgttttaaatccctgtaacagactcgctcagggtgaacttatgttctctgtgc
1 V E Q C F Y I G N F L F K Y P C N R L V R V E L M F L C
4943 aatgtcaacaaaaatttcttcaatcggttcgacctaa 4908
29 N V N K N F F N R S T *
182ORF070
1031 gtgatgggttcggctccacaaaaccagaaacttgcgtcgtgagtaaaactagaatatctttcaattctagtacttcgccaattgcgt
1 V M V R L H Q N Q K L R L S K L E Y L S I L V L R Q L R
947 tacgtaacggaattgaagccccgttgcgttgcattga 912
29 Y V T E L K P R L W H *
182ORF071
11741 atgggtttgcattatgggtgccacaaggcgctcaaaagggttaagggaattttctttaatgatactcgcaattacaatcggtttg
1 M V L H Y G C H K A L K V V K E F S L M I L A I T I V L
11825 actttgatttgcgttgcgttaactgtactttaa 11857
29 T L I C L F V T V L *
182ORF072
11723 atgtttacattaaatgccgtcattgtttcaaaactttaatgtcgttttctccgatcctaagaaagtaactacaggtacatcacgt
1 M F T L N A V I V S N F N V V S P D P K K V T T G T S R
11639 ttcaattcaatggtgttagcaaaagcgataa 11610
29 F N S M V L A K R *
182ORF073
2876 gtgaagccgcctttgtatgctttacgtaagtctttatcaaaccttaagacaaaataggaaaccattgtttgaaagttgatttt
1 V K P P L Y A L R K S L S N P K D K I G N H C L K V D F
2792 ccattgttagcttttagccaatctttgtaa 2763
29 P C V A F S Q S L *
182ORF074
8923 gtgattgataaattttgtttcaaatctgctcgttttgcgtcataaaacggataatcaaaatcaacaattgttttcggcc
1 V I D K F C F K F C S F F V I K R I I K I K Q L F S A
8839 aacttcaatcggttcttttcgagataa 8813
29 N F N T F F S R *
182ORF075
7463 gtgttacattatctggaatattttcgatatctgccactttacctgccaagaggttcaaaccgtttttcttttcagaaacatagt
1 V L H Y L E Y F R Y L P L Y L P R G S N R F L F Q K H S
7379 tgtttacttgttgcctgctcccatga 7353
29 C L L V V L L P *
182ORF076
2426 atgagtgtggagaatgttcgatcttcttttgccttttacaccatttgaaaccatttttgataaacatgaaagcataaactct

328

1 M S V E N V R S S F A S L H H L K P F L N N H E S I N S
2342 ccgtcaaatttttcgttgggaaataa 2316
29 P S N F S L W K *
182ORF077
11858 atgaaggaacgtatgttgttgtgctagaggttagaggggttacatttgaaaattgtctattctctaataatctctcaagcaatta
1 M K E R M L L L L E V E G L H L K I V Y S L I S L K Q L
11942 tcaaaacagcttttccgatgtaa 11965
29 S K Q L F P M *
182ORF078
7671 gtgcctacaatatattgggttcttttaatttaataatgaaattccatgcttttcttgttgaagtttggtagctactcgattgctc
1 V P T I F G S F N L M K F H A F L V C K F G V A T R L L
7587 ttgtgccatacattgagaagtaa 7564
29 F V P Y I E K *
182ORF079
7488 gtgaaagataagtttgatccaagctgtgttacattatctggaatattttcgatatctgccactttacctgccaagaggttcaaa
1 V K D K F D P S C V T L S G I F S I S A T L P A K R F K
7404 ccgttttcttttccagaaacatag 7381
29 P F S F S E T *
182ORF080
4473 gtgtgctatttgcgtgacaaagcttcagttgttgcctccgtagctcttctcgcaatgcttcaagatgttctacaatctttgatc
1 V C Y L L M S K L Q L L L R S L L A M L Q D V L Q S L I
4389 ttgcttcaccgtctgtga 4372
29 L L H R L *

Table 24

Sequence similarities phage 182 and public databases

Phage: 182

Database: nr

Query= sid|110156|lan|182ORF001 Phage 182 ORF|5966-7780|2
(604 letters)

gi 138124 sp P07534 VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) >...	384	e-105
gi 138123 sp P04331 VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9) >...	374	e-103
gi 1429238 gnl PID e1173412 (X99260) tail protein [Bacteriophag...	346	3e-94
gi 215339 (M12456) p9 tail protein (Bacteriophage phi-29) >gi 2...	208	8e-53
gi 1181970 gnl PID e221269 (Z47794) tail protein [Bacteriophage...	62	8e-09
gi 1181968 gnl PID e221267 (Z47794) tail protein [Bacteriophage...	56	6e-07
gi 2500030 sp Q59968 CARA_SULSO CARBAMOYL-PHOSPHATE SYNTHASE SM...	49	8e-05

Query= sid|110157|lan|182ORF002 Phage 182 ORF|2152-3873|1
(573 letters)

gi 118848 sp P19894 DPOL_BPM2 DNA POLYMERASE >gi 76896 pir JQ0...	665	0.0
gi 1429230 gnl PID e1173404 (X99260) DNA polymerase [Bacterioph...	657	0.0
gi 118849 sp P03680 DPOL_BPPH2 DNA POLYMERASE (EARLY PROTEIN GP...	654	0.0
gi 118851 sp P06950 DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP...	654	0.0
gi 15732 (X53371) DNA polymerase (AA 1-575) [Bacteriophage phi-29]	651	0.0
gi 15734 (X53370) DNA polymerase (AA 1-575) [Bacteriophage phi-29]	651	0.0
gi 1572479 gnl PID e242301 (X96987) DNA polymerase [Bacterioph...	565	e-160
gi 1072656 pir S51275 DNA polymerase - phage CP-1 >gi 836593 g...	301	1e-80
gi 118847 sp P22374 DPOM_ASCIM PROBABLE DNA POLYMERASE >gi 8385...	71	3e-11
gi 461962 sp P33537 DPOM_NEUCR PROBABLE DNA POLYMERASE >gi 2833...	65	1e-09
gi 461963 sp P33538 DPOM_NEUTIN PROBABLE DNA POLYMERASE >gi 1018...	62	1e-08
gi 1084487 pir S41618 DNA polymerase - slime mold (Physarum po...	61	3e-08
gi 2435429 (AF012250) unassigned reading frame (possible DNA po...	61	3e-08
gi 578157 gnl PID e246743 (X52106) DNA polymerase [Neurospora i...	59	1e-07
gi 2147969 pir S72369 probable DNA-polymerase - Gelasinospora ...	58	2e-07
gi 2147968 pir S62752 probable DNA-polymerase - Gelasinospora ...	58	2e-07
gi 3511140 (AF061244) B type DNA polymerase [Agrocybe aegerita]	57	3e-07
gi 118850 sp P10479 DPOL_BPPRD DNA POLYMERASE (PROTEIN P1) >gi ...	56	6e-07
gi 578144 (X63909) putative DNA-polymerase, B-type [Morchella c...	47	3e-04
gi 232013 sp P30322 DPOM_AGABT PROBABLE DNA POLYMERASE >gi 3208...	46	6e-04

Query= sid|110159|lan|182ORF004 Phage 182 ORF|4626-5954|3
(442 letters)

gi 138117 sp P13849 VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN ...	309	2e-83
gi 138118 sp P07531 VG8_BPPZA MAJOR HEAD PROTEIN (LATE PROTEIN ...	305	3e-82
gi 1429236 gnl PID e1173410 (X99260) major head protein [Bacter...	300	1e-80
gi 1181958 gnl PID e221257 (Z47794) major head protein [Bacteri...	152	6e-36

Query= sid|110160|lan|182ORF005 Phage 182 ORF|12651-13700|3
(349 letters)

gi 137932 sp P15132 VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE PR...	52	8e-06
gi 1429242 gnl PID e1173416 (X99260) morphogenesis protein [Bac...	48	7e-05
gi 137933 sp P07538 VG13_BPPZA MORPHOGENESIS PROTEIN 1 (LATE PR...	47	2e-04

Query= sid|110161|lan|182ORF006 Phage 182 ORF|14995-16026|1
(343 letters)

gi 137944 sp P11014 VG16_BPPH2 ENCAPSIDATION PROTEIN (LATE PROT...	402	e-111
gi 137945 sp P07541 VG16_BPPZA ENCAPSIDATION PROTEIN (LATE PROT...	402	e-111
gi 1429245 gnl PID e1173419 (X99260) encapsidation protein [Bac...	381	e-105
gi 1181972 gnl PID e221271 (Z47794) encapsidation protein [Bact...	159	2e-38

Query= sid|110162|lan|182ORF007 Phage 182 ORF|7795-8775|1
(326 letters)

gi 1429239 gnl PID e1173413 (X99260) upper collar protein [Bact...	271	5e-72
gi 137915 sp P07535 VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR ...	256	1e-67
gi 137914 sp P04332 VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ...	256	2e-67
gi 1181960 gnl PID e221259 (Z47794) connector protein [Bacterio...	148	6e-35

Query= sid|110163|lan|182ORF008 Phage 182 ORF|14105-14983|2
(292 letters)

gi 4210750 gnl PID e1374037 (AJ132604) LysL protein [Lactococcu...	139	2e-32
gi 462559 sp P34020 LYC_CLOAB AUTOLYTIC LYSOZYME (1,4-BETA-N-AC...	75	8e-13
gi 2327014 (U82823) putative lysozyme [Saccharopolyspora erythr...	64	2e-09
gi 126652 sp P25310 LYCM_STRGL LYSOZYME M1 PRECURSOR (1,4-BETA...	60	2e-08
gi 127789 sp P19386 LYCA_BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	60	2e-08
gi 67761 pir MUBPCP N-acetylmuramoyl-L-alanine amidase (EC 3.5...	59	3e-08
gi 4105636 (AF049087) lys [Leuconostoc oenos bacteriophage 10MC]	59	3e-08
gi 623084 (L02496) muramidase; muramidase [Bacteriophage LL-H]	57	1e-07
gi 127787 sp P15057 LYCA_BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	57	2e-07
gi 126597 sp P00721 LYCH_CHASP N,O-DIACETYLMURAMIDASE (LYSOZYME...	57	2e-07
gi 127788 sp P19385 LYCA_BPCP7 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	57	2e-07
gi 67762 pir MUBPC7 N-acetylmuramoyl-L-alanine amidase (EC 3.5...	56	3e-07
gi 3025168 sp P76421 YEGX_ECOLI HYPOTHETICAL 32.0 KD PROTEIN IN...	53	2e-06
gi 4204413 (AF047001) Lys44 [Oenococcus oeni temperate bacterio...	53	3e-06
gi 2116978 gnl PID d1020940 (D88151) cortical fragment-lytic en...	52	5e-06
gi 2392844 (AF011378) lysin [Bacteriophage sk1]	48	8e-05

Query= sid|110164|lan|182ORF009 Phage 182 ORF|8765-9601|2
(278 letters)

gi 1429240 gnl PID e1173414 (X99260) lower collar protein [Bact...	180	1e-44
gi 137921 sp P04333 VG11_BPPH2 LOWER COLLAR PROTEIN (LATE PROTE...	171	5e-42
gi 215341 (M12456) p11 lower collar protein [Bacteriophage phi-29]	98	9e-20
gi 224162 prf 1011232B protein p11, lower collar [Bacteriophage...	97	1e-19
gi 535260 (Z30339) STARP antigen [Plasmodium reichenowi]	50	1e-05
gi 4049753 (AF063866) ORF MSV230 hypothetical protein [Melanopl...	49	4e-05
gi 2131557 pir S70306 hypothetical protein YEL077c - yeast (Sa...	48	5e-05
gi 131782 sp P12753 RA50_YEAST DNA REPAIR PROTEIN RAD50 (153 KD...	48	7e-05
gi 2131309 pir S70305 hypothetical protein YBL113c - yeast (Sa...	47	2e-04
gi 499325 (Z26314) STARP antigen [Plasmodium falciparum]	46	3e-04
gi 3845171 (AE001391) ribosome releasing factor (OO, TP) [Plasm...	46	3e-04
gi 731903 sp P40434 YIR7_YEAST HYPOTHETICAL 197.5 KD PROTEIN IN...	45	5e-04
gi 1632829 gnl PID e276379 (Y08924) AARP2 protein [Plasmodium f...	45	5e-04
gi 1176490 sp P40889 YJWS_YEAST HYPOTHETICAL 197.6 KD PROTEIN I...	45	5e-04
gi 1077300 pir S51848 hypothetical protein HRD1054 - yeast (Sa...	45	5e-04
gi 2425143 (AF020407) Wima [Dictyostelium discoideum]	45	6e-04
gi 1181961 gnl PID e221260 (Z47794) collar protein [Bacterioph...	45	6e-04
gi 2132657 pir S64819 probable membrane protein YLL067c - yeas...	45	8e-04
gi 2133041 pir S65341 probable membrane protein YPR204w - yeas...	45	8e-04
gi 730275 sp P39793 PBPA_BACSU PENICILLIN-BINDING PROTEINS 1A/1...	45	8e-04

Query= sid|110165|lan|182ORF010 Phage 182 ORF|1310-2155|2
(281 letters)

gi 135604 sp P06812 TERM_BPNF DNA TERMINAL PROTEIN >gi 75815 pi...	69	3e-11
gi 1572478 gnl PID e242334 (X96987) terminal protein [Bacteriop...	65	3e-10
gi 1429231 gnl PID e1173405 (X99260) terminal protein [Bacterio...	64	1e-09

Query= sid|110166|lan|182ORF011 Phage 182 ORF|9607-10158|1
(183 letters)

gi 137928 sp P07537 VG12_BPPZA PRE-NECK APPENDAGE PROTEIN (LATE...	51	6e-06
gi 1429241 gnl PID e1173415 (X99260) pre-neck appendage protein...	51	6e-06
gi 137927 sp P20345 VG12_BPPH2 PRE-NECK APPENDAGE PROTEIN (LATE...	50	1e-05

Query= sid|110169|lan|182ORF014 Phage 182 ORF|13716-14108|3
(130 letters)

gi 137936 sp P11188 VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14...	97	6e-20
gi 137938 sp P07539 VG14_BPPZA LYSIS PROTEIN (LATE PROTEIN GP14...	96	8e-20
gi 1429243 gnl PID e1173417 (X99260) lysis protein [Bacterioph...	96	8e-20
gi 215332 (M14782) lysis protein [Bacteriophage phi-29]	94	5e-19

Query= sid|110170|lan|182ORF015 Phage 182 ORF|854-1225|2
(123 letters)

331

gi 15670 (V01155) reading frame 10 (may be gene 4) [Bacterioph...	70	5e-12
gi 138072 sp P06953 VG5A_BPPZA EARLY PROTEIN GP5A >gi 75836 pir...	69	7e-12

Query= sid|110174|lan|182ORF019 Phage 182 ORF|4323-4613|3
(96 letters)

gi 1429235 gnl PID e1173409 (X99260) head morphogenesis protein...	61	2e-09
gi 138111 sp P13848 VG7_BPPH2 HEAD MORPHOGENESIS PROTEIN (LATE ...	57	3e-08
gi 138112 sp P07533 VG7_BPPZA HEAD MORPHOGENESIS PROTEIN (LATE ...	54	1e-07

Query= sid|110180|lan|182ORF025 Phage 182 ORF|548-814|2
(88 letters)

gi 138099 sp P06955 VG6_BPPZA EARLY PROTEIN GP6 >gi 75841 pir ...	55	7e-08
gi 138098 sp P03685 VG6_BPPH2 EARLY PROTEIN GP6 >gi 75840 pir ...	54	2e-07
gi 1429234 gnl PID e1173408 (X99260) gene 6 product [Bacterioph...	54	2e-07

Table 25

Homologies between 182 ORFs and proteins in public databases

Phage: 182

Database: Swissprot

Query= sid|110156|lan|182ORF001 Phage 182 ORF|5966-7780|2
(604 letters)

gi 138124 sp P07534 VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9)	384	e-106
gi 138123 sp P04331 VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9)	374	e-103
gi 2500030 sp Q59968 CARA_SULSO CARBAMOYL-PHOSPHATE SYNTHASE SM...	49	2e-05

Query= sid|110157|lan|182ORF002 Phage 182 ORF|2152-3873|1
(573 letters)

gi 118848 sp P19894 DPOL_BPM2 DNA POLYMERASE	665	0.0
gi 118849 sp P03680 DPOL_BPPH2 DNA POLYMERASE (EARLY PROTEIN GP2)	654	0.0
gi 118851 sp P06950 DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP2)	654	0.0
gi 118847 sp P22374 DPOM_ASCIM PROBABLE DNA POLYMERASE	71	7e-12
gi 461962 sp P33537 DPOM_NEUCR PROBABLE DNA POLYMERASE	65	3e-10
gi 461963 sp P33538 DPOM_NEUIN PROBABLE DNA POLYMERASE	62	3e-09
gi 118850 sp P10479 DPOL_BPPRD DNA POLYMERASE (PROTEIN P1)	56	2e-07
gi 232013 sp P30322 DPOM_AGABT PROBABLE DNA POLYMERASE	46	2e-04
gi 118887 sp P10582 DPOM_MAIZE DNA POLYMERASE (S-1 DNA ORF 3)	46	2e-04

Query= sid|110159|lan|182ORF004 Phage 182 ORF|4626-5954|3
(442 letters)

gi 138117 sp P13849 VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN ...	309	6e-84
gi 138118 sp P07531 VG8_BPPZA MAJOR HEAD PROTEIN (LATE PROTEIN ...	305	7e-83

Query= sid|110160|lan|182ORF005 Phage 182 ORF|12651-13700|3
(349 letters)

gi 137932 sp P15132 VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE PR...	52	2e-06
gi 137933 sp P07538 VG13_BPPZA MORPHOGENESIS PROTEIN 1 (LATE PR...	47	6e-05

Query= sid|110161|lan|182ORF006 Phage 182 ORF|14995-16026|1
(343 letters)

gi 137945 sp P07541 VG16_BPPZA ENCAPSIDATION PROTEIN (LATE PROT...	402	e-112
gi 137944 sp P11014 VG16_BPPH2 ENCAPSIDATION PROTEIN (LATE PROT...	402	e-112

Query= sid|110162|lan|182ORF007 Phage 182 ORF|7795-8775|1
(326 letters)

gi 137915 sp P07535 VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR ...	256	3e-68
gi 137914 sp P04332 VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ...	256	5e-68

Query= sid|110163|lan|182ORF008 Phage 182 ORF|14105-14983|2
(292 letters)

gi 462559 sp P34020 LYC_CLOAB AUTOLYTIC LYSOZYME (1,4-BETA-N-AC...	75	2e-13
gi 126652 sp P25310 LYCM_STRGL LYSOZYME M1 PRECURSOR (1,4-BETA-...	60	5e-09
gi 127789 sp P19386 LYCA_BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	60	5e-09
gi 127787 sp P15057 LYCA_BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	57	4e-08
gi 126597 sp P00721 LYCH_CHASP N,O-DIACETYLMURAMIDASE (LYSOZYME...	57	4e-08
gi 127788 sp P19385 LYCA_BPCP7 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	57	5e-08
gi 3025168 sp P76421 YEGX_ECOLI HYPOTHETICAL 32.0 KD PROTEIN IN...	53	5e-07

Query= sid|110164|lan|182ORF009 Phage 182 ORF|8765-9601|2
(278 letters)

gi 137921 sp P04333 VG11_BPPH2 LOWER COLLAR PROTEIN (LATE PROTE...	171	1e-42
gi 131782 sp P12753 RA50 YEAST DNA REPAIR PROTEIN RAD50 (153 KD...	48	2e-05
gi 1176490 sp P40889 YJW5 YEAST HYPOTHETICAL 197.6 KD PROTEIN I...	45	1e-04
gi 731903 sp P40434 YIR7 YEAST HYPOTHETICAL 197.5 KD PROTEIN IN...	45	1e-04
gi 730275 sp P39793 PBPA_BACSU PENICILLIN-BINDING PROTEINS 1A/1...	45	2e-04
gi 1168610 sp P41696 AZF1 YEAST ASPARAGINE-RICH ZINC FINGER PRO...	44	3e-04

333

gi 731587 sp P38900 YH19_YEAST HYPOTHETICAL 70.1 KD PROTEIN IN ...	44	3e-04
Query= sid 110165 lan 182ORF010 Phage 182 ORF 1310-2155 2 (281 letters)		
gi 135604 sp P06812 TERM_BPNF DNA TERMINAL PROTEIN	69	8e-12
Query= sid 110166 lan 182ORF011 Phage 182 ORF 9607-10158 1 (183 letters)		
gi 137928 sp P07537 VG12_BPPZA PRE-NECK APPENDAGE PROTEIN (LATE...	51	2e-06
gi 137927 sp P20345 VG12_BPPH2 PRE-NECK APPENDAGE PROTEIN (LATE...	50	3e-06
Query= sid 110169 lan 182ORF014 Phage 182 ORF 13716-14108 3 (130 letters)		
gi 137936 sp P11188 VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14)	97	2e-20
gi 137938 sp P07539 VG14_BPPZA LYSIS PROTEIN (LATE PROTEIN GP14)	96	2e-20
Query= sid 110170 lan 182ORF015 Phage 182 ORF 854-1225 2 (123 letters)		
gi 138072 sp P06953 VG5A_BPPZA EARLY PROTEIN GP5A	69	2e-12
Query= sid 110174 lan 182ORF019 Phage 182 ORF 4323-4613 3 (96 letters)		
gi 138111 sp P13848 VG7_BPPH2 HEAD MORPHOGENESIS PROTEIN (LATE ...	57	9e-09
gi 138112 sp P07533 VG7_BPPZA HEAD MORPHOGENESIS PROTEIN (LATE ...	54	4e-08
Query= sid 110180 lan 182ORF025 Phage 182 ORF 548-814 2 (88 letters)		
gi 138099 sp P06955 VG6_BPPZA EARLY PROTEIN GP6	55	2e-08
gi 138098 sp P03685 VG6_BPPH2 EARLY PROTEIN GP6	54	5e-08

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BLASTP 2.0.8 [Jan-05-1999]

Query= sid|110156|lan|182ORF001 Phage 182 ORF|5966-7780|2
(604 letters)

>gi|138124|sp|P07534|VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9)
>gi|75849|pir||WMBP9Z gene 9 protein - phage PZA
>gi|216058 (M11813) tail protein [Bacteriophage PZA]
Length = 599

Score = 384 bits (975), Expect = e-105
Identities = 231/610 (37%), Positives = 344/610 (55%), Gaps = 36/610 (5%)

Query: 6 TNVLLANVPFDNTYTHTRWFKTQQEQESYFNSFPVLNENRDCSYQRDTQLGGVFRVDKH 65
TNV++LA+VPF N Y +TRWF + Q ++FNS + E ++Q + V
Sbjct: 9 TNVRILADVPPSNDYKNTWRFTSSSNQYNWFNSKTRVYEMSKVTFQGFRENKSYISVSLR 68

Query: 66 KDALYACNYLIFKNESETYPSKWQYAFVTDIEYKNDNTSFVTFEIDVLQTYRFDIGIRESF 125
D LY +Y++F+N + Y +KW YAFVT++EYKN T++V FEIDVLQT+ F+I +ESF
Sbjct: 69 LDLLYNASYIMFQNAD-YGNKWFYAFVTELEYKNVGTITYVHFEIDVLQTMFNIKFQESF 127

Query: 126 IAKEHPQLYYSNGIPFINTIEESLDYGREYTTTNTVTFHPNDGVNFLVILTSEAM--PVG 183
I +EH +L+ +G P INTI+E L+YG EY +V P D + FLV+++ M G
Sbjct: 128 IVRHVKLWDDGTPTINTIDEGLNYGSEYDIVSVENHRPDDMMFLVVISKSIMHGTA 187

Query: 184 DKEDKSG---GSIVGGSPSPSYLLPINSSGEVYKPN-GAGNANFGEYMAFLT---TKEP 236
+ E + S+ G P P YY+ P G+V K G NAN + LT +++
Sbjct: 188 EAESRLNDINASLNGMPQLCYIHPFYKDGKVPKTFIGDNNANLSPIVNMLTNIFSQKS 247

Query: 237 FLNKIVGMVYTSYTGIPFIVDHANKTVRYNAGGSYKIMLPTYASDPTGMTKTFAPFCVKE 296
+N IV MYVT Y G+ + +K ++ + + + A D G + T VK+
Sbjct: 248 AVNNIVNMYVTIDYIGLKLKYNGDKELKLDKDMFEQAGI---ADKXGNVDTIF---VKK 301

Query: 297 ARTFVPKRIDLVGNVYNYFREAFPFNVKESKLFMYPYCLIEITDTKGHVMTLRPEYLTGG 356
+ ID G+ + F + +ESKL MYPYC+ E+TD KG+ M L+ EY+
Sbjct: 302 IPDYETLEID-TGDKWGGFTKD-----QESKLMYPYCVTEVDFKGNHMLKTEYIDNN 355

Query: 357 KLSVYVKGSLGISNKMVIEPIDYDVSNSTI----ITNLSDKMLIDNDPNDVGVS DYASA 412
KL + V+GSLG+SNKV DY+ S +T D LI+N+PND+ + +DY SA
Sbjct: 356 KLKIQVRGSLGVSNKVAYSIQDYNAGGSLSGGDRLTASLDTSLINNNPNDAIINDYLSA 415

Query: 413 FMQGNKNSLIAQEQNIRNTFRHGMGNSAMSTGGAIFSALESNNPFVGLTNIMGAGQQVNN 472
++QGNKNSL Q+ +I GM +S G ++ +PF +++ G N
Sbjct: 416 YLQGNKNSLENQKSSILFNGIVGMLGGGVSA---ASAVGRSPFGLASSVTGMTSTAGN 471

Query: 473 YVSEKENGLNLLAGKVADIENIPDNVTQLGSNLSFTTGN-FQNYVQLRFKQIKYAYATRL 531
V + + L K ADI NIP +T++G N +F GN ++ Y ++ KQ+K EY L
Sbjct: 472 AVL D---MQALQAKQADIANIPPQLTKMGNTAFDYGNGYRGVYVIK-KQLKAEYRRSL 526

Query: 532 DRYFSMYGTSKNRVATPNLQTRKAWNFILKEPNIVGTMSNDVLTRVKQIFSAGVTLWHT 591
+F YG K NRV PNL+TRKA+N+I+ K+ I G ++N+ L ++ IF G+TLWHT
Sbjct: 527 SSFPHKYGYKINRVKKPNLRTKAYNYIQTDCPISGDINNNDLQEIRTI FDNGITLWHT 586

Query: 592 NDVLNYNQDN 601
+D+ NY+ +N
Sbjct: 587 DDIGNYSVEN 596

Query= sid|110157|lan|182ORF002 Phage 182 ORF|2152-3873|1
(573 letters)

>gi|118848|sp|P19894|DPOL_BPM2 DNA POLYMERASE >gi|76896|pir||JQ0161
DNA-directed DNA polymerase (EC 2.7.7.7) - phage M2
>gi|215509 (M33144) DNA polymerase [Bacteriophage M2]
Length = 572

Score = 665 bits (1697), Expect = 0.0
Identities = 327/589 (55%), Positives = 420/589 (70%), Gaps = 38/589 (6%)

Query: 3 KKYTGDFETTTDLNDCRVWSWGVCDIDNVNMTFGLIEDSFFEWCKMQGSTDIYFHNEKF 62
K ++ DFETTT L+DCRVW++G +I N+DN G +D F +W M+ D+YFHN KF

335

Sbjct: 4 KMFSCDFETTTKLDDCRVWAYGYMEIGNLDNYKIGNSLDEFMQWV-MEIQADLYFHNLP 62

Query: 63 DGEFMSLWLFKNGFKWCKEAKEDRTFSTLISNMGQWYALEICWEVNYXXXXXXXXXXXXX 122
DG F+++WL ++GFKW E + T++T+IS MGQWY ++IC+

Sbjct: 63 DGAFIVNWLEQHGFKWSNEGLPN-TYNTIISKMGQWYMIDICFGYK-----GKRKL 112

Query: 123 XXIIYDSLKKYPPFPVKQIAEAFNFIKKGEIDYTKERPIGYKPTKDEWEYLKNDIQIMAM 182
+IYDSLKK PFPVK+IA+ F P+ KG+IDY ERP+G++ T +E+EY+KNDI+I+A

Sbjct: 113 HTVIYDSLKKLPFPVKKIADQFLPLLKGDIDYHTERPVGHEITPEEYIYKNDIEIAR 172

Query: 183 ALKIQFDQGLTRMTRGSDALGDYKDWLKGKSTFKQWFPILSLGFDKDLRKAYKGGFT 242
AL IQF QGL RMT GSD+L +KD L F + FP LSL DK++RKAY+GGFT

Sbjct: 173 ALDIQFKQGLDRMTAGSDSLKGFKDILST----KFKNVFPLSLPMDKEIRKAYRGGFT 228

Query: 243 WVNKVFQKGEIGDGVFDVNSLYPSQMYRPLPYGTPLFYEGEYKPNNDYPLYIQNIKVR 302
W+N ++ KEIG+G+VFDVNSLYPSQMY RPLPYG P+ ++G+Y+ + YPLYIQ I+

Sbjct: 229 WLNDKYEKEIGEGMVFDVNSLYPSQMYSRPLPYGAPIVFGQKYEKDEQYPLYIQRIRFE 288

Query: 303 FRLKEGYIPTIQVKQSSLFIQNEYLESSVNKLGVDELIDLTLTNVDLELFFEHYDILEIH 362
F LKEGYIPTIQ+K++ F NEYL++S GV E ++L LTNVDLEL EHY++ +

Sbjct: 289 FELKEGYIPTIQIKKNPFFKGYEYLKNS----GV-EPVELYLTNVDLELIQEHYELYNVE 343

Query: 363 YTYGYMFKASCDMFKGWIDKWEVKNTEGARKANAKGMLNSLYGKFGTNPDITGKVPYM 422
Y G+ F+ +FK +IDKW VK EGA+K AK MLNSLYGKF +NPD+TGKVPY+

Sbjct: 344 YIDGFKFREKTGLFIDKIDKWTYVKTHEEGAKQLAKMLNSLYGKFASNPDPVTGKVPYL 403

Query: 423 GEDGIVRLTLGEEELRDPVYVPLASFTWANGRYTTITTAQKCFDRIIYCDTDSIHLVGE 482
+DG + +G+EE +DPVY P+ P+TAW R+TTIT AQ C+DRIIYCDTDSIHL GTE

Sbjct: 404 KDDGSLGFRVGDDEYKDPVYTPMGVFITAWARFTTITAAQACYDRIIYCDTDSIHLTGTE 463

Query: 483 VPEAIDHLVDPKGLGYWGHSTFQRAKFIQKT-----YVEEIDGEL----- 524
VPE I +VDPKGLGYW HESTF+RAK++RQKT YV+E+DG+L

Sbjct: 464 VPEIIDKIDVDPKGLGYWGHSTFQRAKYLQKTYIQDIYVKEVDGKLKECSPDEATTTKF 523

Query: 525 NVKAGMPDRIKEIVTFDNFEVGFSSYKLLPKRTQGGVVLVDTMFTIK 573
+VKAGM D IK+ VTFDNF VGFSS GK P + GGVLVD++FTIK

Sbjct: 524 SVKAGMTDTIKKVTDFDNFAVGFSMGPFPVQVNGGVVLVDVFTIK 572

Query= sid|110159|lan|182ORF004 Phage 182 ORF|4626-5954|3
(442 letters)

>gi|138117|sp|P13849|VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN GP8)
>gi|75845|pir||WMBP89 gene 8 protein - phage phi-29
>gi|215325 (M14782) major head protein [Bacteriophage
phi-29] >gi|225362|prf||1301270B gene 8 [Bacillus sp.]
Length = 448

Score = 309 bits (783), Expect = 2e-83
Identities = 176/440 (40%), Positives = 250/440 (56%), Gaps = 27/440 (6%)

Query: 4 KITEQDVLRTNVETPVQLMTAIYNSSSSLFQANVMPNADNIEAVGAGITRLDVVKNF 63
+IT DV + + ++ AI NS F++ VP+ A+N+ VGAGI V+N+F

Sbjct: 2 RITFNDVKTSLGITESYDIVNAIRNSQGDNFKSYVPLATANNVAEVGAGILINQTVQND 61

Query: 64 ISTLVDRIGKVIRYKSWRNPLKMFKGNMPLGRITIEEIFVDIAQEHKFNPDSESVTVFK 123
I++LVDRIG VVIR S NPLK FKKG +PLGRITIEEI+ DI +E +++ +E+ VF+

Sbjct: 62 ITSLVDRIGLVVIRQVSLNNPLKFKKGQIPLGRITIEEIYTDITKEQYDAEEAEQKVFE 121

Query: 124 QEVDPVKTLFHEINREGYKQTIQEAWLEKFTSWDNFNSFVAGVMNALYTGDEVSEFEY 183
+E+P+VKTLPHE NR+G+Y QTIQ+ L+ AF SW NF SPV+ ++NA+Y EV E+EY

Sbjct: 122 REMPNVKTLPHERNRQGFYHQTIQDDSLKTAFPVSWGNFESFVSSIINAIYNSAEVDEY 181

Query: 184 TKLLIANVQEKELFKEIEIGEITESNA--KEFIRKIKSTSNKLEFM--SSAYNAQGVKTS 239
KLL+ NY K LP ++I E T S EF++K+++T+ KL S +N+ V+T

Sbjct: 182 MKLLVDNYYSKGLFTTVKIDEPTSSGTALTEFVKMRATARKLLTPQGSRDWNSMAVRTR 241

Query: 240 TSKSDQYXXXXXXXXXXXXXXXXXFNMSKTDVFGHKIVIDEFPKKEGESSNIVAVIV 299
+ D + FNM++TDF+G+ VID F S+ + AV+V

Sbjct: 242 SYMEDLHLIIDADLEAELDVDVLAKAFNMNRDFTLGNVTVIDGF-----ASTGLEAVLV 295

Query: 300 DSEWFMIYDKLYKTSLYNPEGLYWNYNLHHHQLYSTSQFGNAFAVKSATKPVTKVAF 359
D +WFM+YD L+K ++ NP GLYWN+ H Q S S+F NAVAFAV VT+V +

Sbjct: 296 DKDWFMYVDNLHKMETVRNPRGLYWNYYYHVWQTLVSRSFANAVAFVSGDVPVAVTQVIVS 355

Query: 360 SATTSVVKGSSKDIALTFTPEATNQGEVSSAPALVKATVKTAGKATAVTVEGLEV 419

336

+V +G + V ATN + V V G +T + G
 Sbjct: 356 PNIAAVKQGGQQQFT---AYVRATNAKDHKV-----VWSVEGGSTGTAI---TG 398

Query: 420 QSLVTFATAIGGQQATVLVTV 439

L++ + Q TV TV

Sbjct: 399 DGLLSVSGNEDNQLTVKATV 418

Query= sid|110160|lan|182ORF005 Phage 182 ORF|12651-13700|3
 (349 letters)

>gi|137932|sp|P15132|VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE
 PROTEIN GP13) >gi|75858|pir||WMBP23 gene 13 protein -
 phage phi-29 >gi|215331 (M14782) morphogenesis protein
 [Bacteriophage phi-29] >gi|225368|prf||1301270H gene 13
 [Bacteriophage phi-29]
 Length = 365

Score = 51.5 bits (121), Expect = 8e-06

Identities = 44/166 (26%), Positives = 70/166 (41%), Gaps = 14/166 (8%)

Query: 6 NEQIARGQTIKILSKYGYNKNSQVGVVANLHWESA---GLNPNSNEXXXXXXXX-QWT 61

+E Q I LS G+ K + G++ N+ ES GL N +E QWT

Sbjct: 12 SEMKVNAQYILNLYLSSNGWTKQAICGMLGNMQSESTINPGLWQNLDEGNTSLGFLVQWT 71

Query: 62 PKSNLRYRQAQICGLSNAKAETLEGQAEIIAQGDKTGQWMDNTPVSSAGYTNPQTLSAFKQ 121

P SN A GL ++ II + + QW++ ++ Y K

Sbjct: 72 PASNYINWANSQGLPYKMDMS--ELKRIIWEVNNNAQWINLRDMTFKEY-----IKS 121

Query: 122 SANIDVATINFMCHWERPGKLHIEERLDAQAYSKHIDSGGGGVK 167

+ + F+ +ERP + ER D A+ + K++ G GGGG++

Sbjct: 122 TKTPRELAMIFLASYERPANPNQPERGDQAEYWKNLGGGGGGLQ 167

Query= sid|110161|lan|182ORF006 Phage 182 ORF|14995-16026|1
 (343 letters)

>gi|137945|sp|P07541|VG16_BPPZA ENCAPSIDATION PROTEIN (LATE PROTEIN
 GP16) >gi|75861|pir||WMBP16 gene 16 protein - phage PZA
 >gi|216065 (M11813) morphogenesis protein C
 [Bacteriophage PZA]
 Length = 332

Score = 402 bits (1023), Expect = e-111

Identities = 186/332 (56%), Positives = 244/332 (73%), Gaps = 2/332 (0%)

Query: 11 EKNLYNPNNALGFNCLMLFVIGARGIGKTYGYKFFVNRFIKHGEQFIYLRPFKTELKK 70

+K+L+YNP L ++ ++ FVIGARGIGK+Y K + +NRFIK+GEQFIY+RR+K EL K

Sbjct: 2 DKSLFYNPQKMLSYDRILNFVIGARGIGKSYAMKVYPINRFIKYGEQFIYVRRYKPELAK 61

Query: 71 IPQFFKTMKEFPDHLKLEVKGEFYCDDKLMGWAVPLSTWIEKSNEYPEVRTILFDEF 130

+ +F +A+EFPDHL VKG+ FY D KL GWA+PLS W EKS N YP V TI+DEF+

Sbjct: 62 VSNYFNDVAQEFDPDHELTVKGRRFYIDGKLAGWAIPLSVWQSEKSNAYPNVSTIVDEFI 121

Query: 131 IEKSKITYLPNEAEALLNMETVFRRTNTRCVMLSNATSVVNPYFLYFNLQPDNLKRPN 190

EK Y+PNE ALLN+M+TVFR R RC+ LSNA SVVNPYFL+FNL PD+NKRPN

Sbjct: 122 REKDNSNYIPNEVSALLNMDTVFRNRERVRCICLSNAVSVVNPYFLFNLVDPVNRPN 181

Query: 191 LYQDRGILIELCDSKDFAEVKRETPFGRRLIRGTEYEDFSINNEFVNDSDTFIEKRSKNSS 250

+Y D LIE+ DS DF+ +R+T FGRLI GTEY + S++N+F+ DS FIEKRSK+S

Sbjct: 182 VYDD--ALIEIPDSLDFSSERRKTRFGRLIDGTEYGEMSLDNQFIGDHSVFIKRSKDSK 239

Query: 251 FLCAIAFEGKIFGYWIDAETGCVVVSVDYQPNNTNHFYAMTTKDHEENRLLMKNNRNNYYL 310

F+ +I + G G W+D G +YV + P+T + Y +TT D EN +L+ N++NNY+L

Sbjct: 240 FVFSIVYNGFTLGWVDVNVQGLMYVDTAHPSTKNVYTLTDDLNNENMLITNYKNNYHL 299

Query: 311 STVAKAPKNSYLRFDNIVIKNLHYDLFNKMKI 342

+A AF N YLRFDN VI+N+ Y+LF KM+I

Sbjct: 300 RKLASAFMNGYLRFDNQVIRNIAIELFRKMRI 331

Query= sid|110162|lan|182ORF007 Phage 182 ORF|7795-8775|1
 (326 letters)

>gi|1429239|emb|CAA67658| (X99260) upper collar protein
 [Bacteriophage B103]

337

Length = 308

Score = 271 bits (685), Expect = 6e-72
Identities = 131/275 (47%), Positives = 187/275 (67%), Gaps = 5/275 (1%)

Query: 36 YYEYRRQLTLLTFQLFEWENLPKSIDPRYLEIALHTNGYLGFCKDPTLGFMVCAGAEDG 95
+Y HY + L L +QLFEWE LP S+DP YLE ++H GY+GF+KDP +G++ C GA G
Sbjct: 22 WYHYHYQLCSLAYQLFEWERLPPSVDPFSYLEKSIHQFGYGVGYKDPRIIGYIACQGALSG 81

Query: 96 QIDHYHNPFIPTANEAMYHKRYPVLRYYDDDDKSKCIMLYNNDLKVPTLPSLHRFALDMA 155
+DHY+ P F A+ Y + + Y D +K+ + +YNNDLK TLP+L FA D+A
Sbjct: 82 TVDHYNLPDRFHFASSVGYQNTFKLYNSDMKEKNMGVAIYNNDLKCSLTLPALMFQAQDLA 141

Query: 156 DINQISRVNRRQAQKTPVVIQTDKQYFSLQAYNQIDENNQAVFVDKMEFDESFNWQT 215
++ +I VN+ AQKTPV+I ++ SL YNQ + N +FV + ++ D + V++T
Sbjct: 142 ELKEIIAVNQNAQKTPVLIAANDNNQLSLKNIYNQYEGNAPVIFVHESLDLD-NLKVFKT 200

Query: 216 NAPIYVVDKLRSELNEVWNEVLTFGLINNANVDKTARVQTSEVLSNNEQIESSGNILLKSR 275
+APIYVVDKL ++ N VVNEV+T+LGI NAN++K R+ TSEV SN+EQIESSGNI LK+R
Sbjct: 201 DAPIYVVDKLNQKNNAVWNEVMTYLGIXNANLEKERMVTSEVDSNDEQIESSGNIYLKAR 260

Query: 276 KEFCDRVNRVFGDELQKIDVFRDQVRLQALAA 310
+E C++++ ++G L VKFR D V Q++L A
Sbjct: 261 QEACNKISELYGLNL----KVKFRYDIVEQMRLNA 291

Query= sid|110163|lan|182ORF008 Phage 182 ORF|14105-14983|2
(292 letters)

>gi|4210750|emb|CAA10710| (AJ132604) LysL protein [Lactococcus
lactis]
Length = 235

Score = 139 bits (347), Expect = 2e-32
Identities = 85/210 (40%), Positives = 114/210 (53%), Gaps = 14/210 (6%)

Query: 2 MNGIDISSYQTGIDLSKVPDFVNIKATGGTGYVNPDCDRAQQALSIGKKIGVYHFAHE 61
MNGIDISSYQ ++ VP DFV IKAT GT Y+NP + Q + K +G YHFA
Sbjct: 1 MNGIDISSYQAEINAGIVPSDFVIIKATEGTNYINPTWEEQAGQVIQTNKLLGFYHFAS- 59

Query: 62 RGLEGTQQEQAQFFLDNIKGYIGKAVLILDFEGS--NQKDVNWAFLDYVYNKTGVKAW 119
G P EA FF+ +K YIGKAVL+LDPE N A+ FL+ V KTG+
Sbjct: 60 ---VGNPIAEADFFISVVKYIGKAVLVLDPEAGAINANGVNGARQFLNRVKEKTGINPM 116

Query: 120 FYTYTANLNTIDFSSIAKGDYGLWVAEYGSNQPGYSQPAPPKTNN-----FPIVACQF 174
Y + ++S+I+ + LWVA+Y S P GY + P T+ + A Q+
Sbjct: 117 IYMSSDVTRQFNWSTISSTN-PLWVAQYASMNPTGYQ--SEPWTGKGYGAWSSAAIHQY 173

Query: 175 TSKGRLPGYNGNLDLNVFYGDGNTWDLYVG 204
+S G L ++GNLD+N+ Y + N W C
Sbjct: 174 SSAGSLSNWSGNLDINLAYINANQWKS LAG 203

Query= sid|110164|lan|182ORF009 Phage 182 ORF|8765-9601|2
(278 letters)

>gi|1429240|emb|CAA67659| (X99260) lower collar protein
[Bacteriophage B103]
Length = 293

Score = 180 bits (451), Expect = 1e-44
Identities = 115/296 (38%), Positives = 161/296 (53%), Gaps = 33/296 (11%)

Query: 3 LKRYIESFTYYQPELSRKRIEIVGRKQLFDYDFYDETKRAEFETKFINHFYLRIGSE 62
L YIE ++ Y+ LS E+IE GR +LDFD YP +DE+ R FET FI +FY+REIG E
Sbjct: 8 LSTYIEMWSQYETGLSMAEKIEKGRPKLFDYDFYPIFDESIRKVFETHFIRNFYMRIGFE 67

Query: 63 TMGSPKFNLDYELNLMNPYWNKMFSLNLEEF-PIFDDMDYTIDEKQKLLNEIDTNIKAR 121
T G FKFNL+ +L +NMPY+NK+F S L ++ P+ + T K+ DT NR
Sbjct: 68 TEGLPFKNLETLIINMPYFNKLFESSELIKYPLENTLNTTGNKKN-----DTERNDNR 122

Query: 122 D-----ESKNQTKQVDQTDNRNKNTRDTGTT-----DSFSRNTYTDTPQKDLRIASNG 169
D + K+ TK D+T+ + D TT D+F+R +D P L + +N

338

Sbjct: 123 DTGSMKADGKSNTKTSKTNATGSSKEDGKTGVTDDNFNRKIDSDQPDRLNLTN- 181

Query: 170 DGTGVINYATNITEDLSKETTSSTGVETNNDKTNQNTSRNAS-----EKETKNTD 219
 DG G + YA+ I E+ + ++TG TNN ++ + S S T N

Sbjct: 182 DGQGTLEYASAIENNTNNKRNTTG--TNNVTSSAESESTGSGTSDTVTTDNANTTTNDK 239

Query: 220 INKDQNTKDTITRYKGGKGNTRYADLLEKYRRSVLRIEKMIFREMKEGLFLLVY 275
 +N N +D I GK G YA L++ YR ++LRIEK IF EM + LF+LVY

Sbjct: 240 LNSQINNVEDYIESKIGKSGTQSYASLVQDYRAALLRIEKRFDEMQE--LFMLVY 293

Query= sid|110165|lan|182ORF010 Phage 182 ORF|1310-2155|2
 (281 letters)

>gi|135604|sp|P06812|TERM_BPNF DNA TERMINAL PROTEIN
 >gi|75815|pir|ERBPNP terminal protein - phage NF
 >gi|579177|emb|CAA68440| (Y00363) gene E product (AA
 1-267) [Bacteriophage NF]
 Length = 266

Score = 74.9 bits (181), Expect = 6e-13
 Identities = 73/275 (26%), Positives = 129/275 (46%), Gaps = 37/275 (13%)

Query: 3 VRISKNDRAKLEKIYKSNKARKKYNRLRQK-GVE---ERQLPTVPTSKKRLIDYVKSTN 58
 +RI+ ND+A K+ K+ KA K +R ++K G++ E +LP + + +

Sbjct: 7 IRTNNDKALYAKLV-KNTKA--KISRTKKYKIDLSNEIELPPLESFQ----- 52

Query: 59 MSRSDFNKMLDELVDFAQPYNENYIFEINKRNVAISRAQIKEAQIKTEQAQKAKEEHYKE 118
 +R +FNK + F N+NY F NK + S+A+I E T++AQ+ +E +E

Sbjct: 53 -TREEFNKWKQKQESFTNRANQNYQFVKNKYGIVASKAKINEIAKNTKEAQRIVDEQREE 111

Query: 119 L-----NKVEVKKPTENTIVTPTILTELGADLPFQAIPDFNIDAFPTSPEGVQSYLEN 170
 + K + I++P+ +T G P DFN D S +++ E

Sbjct: 112 IEDKPFISGGKQQTGQRMQILSPSQVT--GISRP----SDFNFDDVRSYARLRTLEEG 165

Query: 171 IG-KQDEQYFDERDQLYYDNFRQAMFTIFNSD--ADDIVRLDLSMGLDLFMKTYVSNFLD 227
 + K Y+D R + NF + + FNSD +D++V L + D F + Y+ F +

Sbjct: 166 MAEKASPDYDRMTQMHNQNFIEIVEKSFNSDWLSDELVERLKKIPDDFFELYLM-FDE 224

Query: 228 MNLDIYDEAEVQKKEQVYSKIAKVIESETGGEV 262
 ++ +Y E E + E + +KI ++ G+V

Sbjct: 225 ISFEYFDSEGEDVEASEAMLNKIHSYLDYRERGDV 259

Query= sid|110166|lan|182ORF011 Phage 182 ORF|9607-10158|1
 (183 letters)

>gi|1429241|emb|CAA67660| (X99260) pre-neck appendage protein
 [Bacteriophage B103]
 Length = 860

Score = 50.8 bits (119), Expect = 6e-06
 Identities = 29/105 (27%), Positives = 56/105 (52%), Gaps = 6/105 (5%)

Query: 8 KRFDGLPAVFKERFSKYPHTEYRYELLDEEVSAIAYLNEVGALVNDMSGYLNIFYEHF 67
 +RF+ L + + + +Y T + + L E+++ +I YLN++G L ND+ N +E

Sbjct: 7 RRFELGEMMVQVYERYLPTAFDESMTLLEKMNKIIEYLNQIGRLTNDVVEBNKVMEMI 66

Query: 68 V-EKLEITNDTLKKWLSGDTLENLINDTVFANYIKEIKRLQILV 111
 + + LE+ +TL+KW +G +L+ I E+K+ + V

Sbjct: 67 LNDGLEDYVKETLEKWEYEGKFADLV-----IQVIDELKQFGVSV 106

Query= sid|110169|lan|182ORF014 Phage 182 ORF|13716-14108|3
 (130 letters)

>gi|137936|sp|P11188|VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14)
 >gi|75860|pir|WMBP29 gene 14 protein - phage phi-29
 >gi|15678|emb|CAA28631| (X04962) gene 14 product (AA

339

1-393) [Bacteriophage phi-29] >gi|225369|prf||1301270J
 gene 14 [Bacteriophage phi-29]
 Length = 131

Score = 96.7 bits (237), Expect = 6e-20
 Identities = 53/131 (40%), Positives = 81/131 (61%), Gaps = 3/131 (2%)

Query: 1 MIEYITQWL-ADDNHLVYGLIIWLMVAMIIDFVLGFTIAKFNKEIDFSSFKAKAGIIVKV 59
 MI ++ +L D+ L+Y L +LMV M++D VLG AK N I FSSFK K G+++KV
 Sbjct: 3 MIAWMQHFLFLETDKLIYWLT-FLMVCMVVDIVLGVLFAKLNPNIKFSSFKIKTGVLIKV 61

Query: 60 AEMVLVVYPIPVAVKFGAVGITMYITMLVGLILSEIYSILGHISDIDDDNNWTDYVKKFL 119
 +EM+L + IP AV F A G+ + T+ L +SEIYSI GH+ +DD +++ + ++ F
 Sbjct: 62 SEMILALLAIPFAVFPFA-GLPLLYTVYTALCVSEIYSIFGHLRLVDDKSDFLLELENFF 120

Query: 120 DGTILNRKDDIK 130
 T + + K
 Sbjct: 121 KRTSGKNKEEK 131

Query= sid|110170|lan|182ORF015 Phage 182 ORF|854-1225|2
 (123 letters)

>gi|15670|emb|CAA24483| (V01155) reading frame 10 (may be gene 4)
 [Bacteriophage phi-29]
 Length = 124

Score = 69.9 bits (168), Expect = 6e-12
 Identities = 39/119 (32%), Positives = 64/119 (53%), Gaps = 3/119 (2%)

Query: 3 IVKSTFDTQTPEGMLQVFNATNGASIPLRNAI-GEVLEKDIILVYSDEVSGFGGAEPSQA 61
 IVK+TFDT+T EG +++FNA G +N G ++E I Y +G A+ +
 Sbjct: 6 IVKATFDTETLEGQIKIFNAQTGGGQSFKNLPDGTII EANAIAQYKQVSDTYGDAK--EE 63

Query: 62 ELVAFPTEDGKTYAGVSAVATKSAKNLIDMMTANPDIKPKISFVEGKSNGGQKFVNLQV 120
 + F DG Y+ +S ++A +LID++T + K+ V+G S+ G F +LQ+
 Sbjct: 64 TVTTIFAADGSLYSAISKTVAEASDLIDLVTTRHKLETFKVKVVGQTSKGNVFFSLQL 122

Query= sid|110174|lan|182ORF019 Phage 182 ORF|4323-4613|3
 (96 letters)

>gi|1429235|emb|CAA67654| (X99260) head morphogenesis protein
 [Bacteriophage B103]
 Length = 101

Score = 60.9 bits (145), Expect = 1e-09
 Identities = 34/96 (35%), Positives = 53/96 (54%), Gaps = 5/96 (5%)

Query: 1 MEIKEHESILNGILESVTGGEARSKIVEHLEALREDYGATTEALTSANSTLEKLLKONEA 60
 ME HE ILN + + + R+++ L+ LR DYG+ + S EKL+ +N
 Sbjct: 3 MERDSHEEILNKLNDPELEHSERTEL---LQQLRADYGSVLSEFSELTSAEKLRAENSD 59

Query: 61 LVISNSKLFRRERAVEPAEN--NEPETDQNTLDDL 94
 L++SNSKLF+ I + E + E + IT++DL
 Sbjct: 60 LIVSNSKLFQVGITKEKEEBIKQEELSETITIEDL 95

Query= sid|110180|lan|182ORF025 Phage 182 ORF|548-814|2
 (88 letters)

>gi|138099|sp|P06955|VG6_BPPZA EARLY PROTEIN GP6
 >gi|75841|pir||ERBP6Z gene 6 protein - phage PZA
 >gi|216047 (M11813) gene 6 product [Bacteriophage PZA]
 >gi|224746|prf||1112171K ORF 6 [Bacteriophage PZA]
 Length = 96

Score = 55.0 bits (130), Expect = 8e-08
 Identities = 28/79 (35%), Positives = 45/79 (56%)

Query: 4 KLMQRNVTSTKVEFSEVIVQDGAPTIVPCEPVVLTGKLSEEKALSAIKRKNPDKNVVVTN 63
K+MQR +T T V +++++ DG + G LS E+A +KRK + V V +
Sbjct: 3 KMMQREITKTTVNVAKMVMVDGEVQVEQLPSETFVGNLSMEQAQWRMKRKYKGEPVQVVS 62

Query: 64 VSHETALYTMPVDKFIELA 82

V T +Y +PV+KF+E+A

Sbjct: 63 VEPNTEVYELPVEKFLEVA 81

Table 26

Secondary structure prediction for ORF 182ORF008

```

1  MMNGIDISSY QTGIDLSKVP CDFVNIKATG GTGYVNPDCD RAFQQALSLG KKIGVYHFAH
   CCCCCCCCCC CCCCCCCCCC CEEEEEEEC CCCCCCCCCC HHHHHHHHHC CCCCEEEEEE
61  ERGLEGTPQQ EAQFFLDNIK GYIGKAVLIL DFEGSNQKDV NWAKAFLDYV YNKTGVKAWF
   CCCCCCCHH HHHHHHHHHC CCCCEEEEEE CCCCCCHH HHHHHHHHHC HCCCCCEEE
121 YTYTANLNTT DFSSIAKGDY GLWVAEYGSN QPQGYSPAP PKTNNFPIVA CFQFTSKGRL
   EEECCCCCCC CCCECCCCC CEEEEEEEC CCCCCCCCCC CCCCCCEEE EEECCCCCCC
181 PGYNGNLDLN VFYGDGNTWD LYVGKKQDQI VPPENKIFDA TSDEFIFTLT TGSTSVFYFD
   CCCCCCCEE EEECCCCCE EEECCCCCC CCCCCCCCCC CCCEEEEEEC CCCCEEEEC
241 GETIFELSDP TQLDHIRGTY NHVHGKEIPS MVWTPEQFDI YLKMYEKKPV YK
   CCEEECCCC CCHHHHCCE CCCCCCEEC CCCCCCHH HHHHCCCC EC

```

Secondary structure prediction for ORF 182ORF014

```

1  MIEYITQWLA DDNHLVYGLI IWLVMAMIID FVLGFTIAKF NKEIDFSSFK AKAGIIVKVA
   CCCCEEECCC CCCCHHHHH HHHHHHHHHC CCCCCHHHH HHHCEEEEEE
61  EMVLVVYFIP VAVKFGAVGI TMYITMLVGL ILSEIYSILG HISDIDDDNN WTDYVKKFLD
   EEEEEEEEC CEECCCEEE EEEEEEEEE EEEEEEEEC CCCCCCCCC CEEEEEEEC
121 GTLNRKDDIK
   CCCCCCEEC

```

Table 27

Enterococcus accession numbers 242/242

gi 2895751 gb AF044978.1 AF044978 [2895751]	gi 4098267 gb U76614.1 BLU76614 [4098267]
gi 4803755 dbj AB026843.1 AB026843 [4803755]	gi 47019 emb Y00116.1 SFAMB1 [47019]
gi 4769001 gb AF140549.1 AF140549 [4769001]	gi 4158179 emb AL035206.1 SC9B5 [4158179]
gi 4760901 gb AF099088.1 AF099088 [4760901]	gi 4165458 emb X79343.1 EF16SSPA [4165458]
gi 4704705 gb AF121254.1 AF121254 [4704705]	gi 4165457 emb X79342.1 EFTRNALA [4165457]
gi 3342117 gb AF076604.1 AF076604 [3342117]	gi 4165456 emb X79341.1 EF23SRNA [4165456]
gi 4688824 emb AJ132470.1 ESP132470 [4688824]	gi 4150978 emb Y14027.1 EFY14027 [4150978]
gi 4732085 gb AF125553.1 AF125553 [4732085]	gi 4127803 emb AJ223161.1 EFAJ3161 [4127803]
gi 4732082 gb AF125552.1 AF125552 [4732082]	gi 2956685 emb Y16413.1 EFENTIJO [2956685]
gi 4732079 gb AF125551.1 AF125551 [4732079]	gi 2665346 emb Y13922.1 EHY13922 [2665346]
gi 4732076 gb AF125550.1 AF125550 [4732076]	gi 4324675 gb AF109375.1 AF109375 [4324675]
gi 4732073 gb AF125548.1 AF125548 [4732073]	gi 4234627 gb AF061013.1 AF061013 [4234627]
gi 4732070 gb AF125547.1 AF125547 [4732070]	gi 4234626 gb AF061012.1 AF061012 [4234626]
gi 4732067 gb AF125546.1 AF125546 [4732067]	gi 4234625 gb AF061011.1 AF061011 [4234625]
gi 4732064 gb AF125545.1 AF125545 [4732064]	gi 4234624 gb AF061010.1 AF061010 [4234624]
gi 4732061 gb AF125544.1 AF125544 [4732061]	gi 4234623 gb AF061009.1 AF061009 [4234623]
gi 4704653 gb AF114715.1 AF114715 [4704653]	gi 4234622 gb AF061008.1 AF061008 [4234622]
gi 4704564 gb AF102550.1 AF102550 [4704564]	gi 4234621 gb AF061007.1 AF061007 [4234621]
gi 4688827 emb AJ238249.1 EFA238249 [4688827]	gi 4234620 gb AF061006.1 AF061006 [4234620]
gi 4680606 gb AF125198.1 AF125198 [4680606]	gi 4234619 gb AF061005.1 AF061005 [4234619]
gi 4633279 gb AF117609.1 AF117609 [4633279]	gi 4234618 gb AF061004.1 AF061004 [4234618]
gi 4633124 gb AF110130.1 AF110130 [4633124]	gi 4234617 gb AF061003.1 AF061003 [4234617]
gi 4590399 gb AF124258.1 AF124258 [4590399]	gi 4234616 gb AF061002.1 AF061002 [4234616]
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gi 4019167 gb U21300.1 CXU21300 [4019167]	gi 3138990 gb AF060241.1 AF060241 [3138990]
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 gi|43333|emb|X16421.1|EFPF54 [43333]
 gi|43331|emb|X62657.1|EFORF3 [43331]
 gi|1065721|emb|X92945.1|EFCAT501 [1065721]
 gi|806551|emb|Z49243.1|EF4110SOD [806551]
 gi|806549|emb|Z49244.1|EF4105SOD [806549]
 gi|505530|emb|X79542.1|EFAS48 [505530]
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 gi|48189|emb|X04388.1|TN1545TR [48189]
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gi|497792|dbj|D31676.1|ENC16RNA9 [497792]
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Table 28

Phage Dp1 complete genome sequence. 56506 nucleotides.

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51591 gatgacctta aaacggaatt tctttatggc caatatgagc ttgtagcatt tcgagactat ttgaaaaaac
51661 ctatttagtca agtattcgtg actgagctcg ttatcaactg cttgactctt tgggtcaatga agattccagc
51731 agtcgtctct atgggagtag gtggaggaaa tcaaatcaat ttactaaaac gacttctcta tagaaatatt
51801 gttctagcac ttgaccttga taacgctggg cagacagcgc aggaaaaact ctaccgacag ttaagcgaa
51871 gcaaggtcgt tagatttttg aactacccta aagagttcta tgataataag tgggataata acgacctcc
51941 ggaattatta aattttaatg atttagtctt gtgaaaattc attttattat gtataataaa gttgaaaat
52011 tttaaaaaga ggtcatatca atatgaaaga agcgaataga ctagtttcta gctatgtag attcgaatgc
52081 tggactgacg aagaatgtat caggaaactt gaactagacc ctgatatgtc aattgctctc gtttattcat
52151 gttatttttg gatgctttat tccatgcaa aaagggttaa atgcttatct cgacatgaca ttgaaagcat
52221 tgcattcgag actatttcaa aatgtttggc aacgttcaaa tcaaaccaag gggccaagt ttcaacttac
52291 cttacaagac tcttcaagaa tagaatagtc ttagaatata ggtacctaaa tgcaccttcc atgaatcgaa
52361 attgtattgt agaagtgacg ttcgatagcg tttcgacaaa tgaagaaggc gacgatttta gtatcctatc
52431 gacagttggc tattgtgaag actacggaaa aattgaaatt gaagcaagtc ttgacttcat gacgcttct
52501 aatacagagt atgcttata ctcgtctgtc attcaaaacg gtccttcagt aagcgacgca gaaattggc
52571 gtgaaatttg agtaagcagg tctgctatta gtcagctcaa gaagtcacta aaaaataaat taaaagattt

52641 tatataactg gtttacaaat cacgtgaatt tcgtgtatat tatatatgaa aggacaaact ttgaaacctt
52711 aaaaacttca aaaatctttc aaccattaaa aacttataaa ggagaatcga tatgggaaaa gtatcaattc
52781 aaaaatcagg aacatttagc tcagggtcta ataacgagtt ttccacactc gctgaccacg gtgacagcgc
52851 aattgtcact ctattgtatg atgacccgga aggcgaagac atggattatt tcgtagtcca cgaagcagac
52921 gttgacggtc gtcgacgcta tatcaattgc aatgctattg gcgaagacgg ggaacacagtc catcctgata
52991 attgtccatt atgcaaaaac ggattccctc gtattgaaaa actatttctt caactttaca accatgatac
53061 gggaaaagtt gaaacatggg accgaggcgg ttcttatgtt caaaagattg ttacatttat caataaatat
53131 ggaagccttg tgaactcagcc ttttgaaatt attcggtcag gagctaaaag tgaccaacga actacttatg
53201 aattccttcc agagcgtccg gaagacagtg ctactcttga agattttcca gaaaagagcg aacttcttgg
53271 aactcctaatt ttgacctcgc acgaagacca aatgtttgac gtggttgacg gcaagttcac tcttcaagaa
53341 gagcggtctt caagtctgtc aaattccagt agaggagcat ctctcgccgc tagacgaggt tccggtcgag
53411 aatcttccca aggtcgaaca gctgaaagaa ctcttcagt tagtcaaga actcctccaa cacgaggtcg
53481 aggattctaa catgagggcg cgagccctct ttattattga ttaagaaaag gaaaataatg gcacaaaaag
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53901 gttagcaata tgacgaagat gcgaattaa atcaaatct ctctgagtt catgaagaa atgcttcaac
53971 ggattgtaga ttcaaggaatt cctgtcatct atcataattc gaaatttgac atgaaatcga tttattggcg
54041 actcggcgtc aaaatgaatg agccagcgtg ggatacatat ttacccgcaa tgcctttaa tgaaaacgag
54111 tctcacagct tgaagagctc tcaactctaa tatgttagga acgaagaaaa cgcagaggtt gcaaaattta
54181 atgacttatt taaaggaatt ccttttagtt taattcctcc tgatgttgc tatatgtat cggcctatga
54251 cctcttgcaa actttcgaa cctatgaatt tcaagaacaa tacttgactc caggaactga acaatgtgaa
54321 gaatataacc tggaaaaagt ctcattgggt cttcataata ttgagatgcc tctaattaaa gttctcttcg
54391 acatggaagt ctacggtgtc gacttagacc aagataagct ggcagaaatt agagaacagt ttaactgcaa
54461 tatgaacgag gctgagcaag agtttcaaca gcttgcagc gaatggcagc ctgaaattga agaacttcga
54531 caaactaatt tccagagcta tcaaaaactc gaaatggatg caagaggtcg agtgacggta agcatttcca
54601 gctcactca attagcaatt ctgttttatg atatcatggg attgaaaagt cctgaaaggg ataaacctag
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54951 gactactctc aacaagaacc tcgttcatgt gcggaattaa gtggcgacga aagtatgca catgcttacg
55021 aacaaaacct ggacctatat cagtttatcg gttcgaaact ttatggtgtt ccctatgaag agtggttaga
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55161 ggtcttatgt acggccgagg ggctaaactc atcgctgagc agatgaatgt atctgtcaaa gaagcgaata
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55301 gcaggacttg ggatatgttc aaacagctac cggtcgaaga agaaggcttc ctgatatgag tcttctgaa
55371 tacgagttcg agtatatcga cgctagcaag aacgaagatt tcgacccctt taactttgac gcagaccaac
55441 agatggacga tactgttctc gaacatatta tgcgaataa ttgggcccag ctgatatagag ctggggatt
55511 taagaagaag caagaatta aagaccaggc aaaagccgaa ggaattctta ttaaggataa cggaggcaag
55581 atagctgatg ctacagccca atgtttgaac tcagttatcc aaggaaacggc agccgacatg actaagtacg
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55721 tgagttacta ggtgaggttc ctatcaagaa cgcaaaacgg ggagcagaaa ggttgacaga agttatgatt
55791 gaagcagcca aggacattat tagtcttcca atgaaatgtg accccagtat agtagaaga tggtaggtg
55861 aagaaattga aatctaaaat ctattcagtt gcatatataa ttctagtagt tattgcaaac cttgtgacaa
55931 tttatttcga acctttaa atgtgaaaggaa ttttaattcc tccaagcagt tgggttatgg gattcacttt
56001 cctgcttata aatctaataa gcaagtacga gaagccaaaa tttgcaggtt ctttgatag ggtagggtta
56071 ttcttacct cgttgatttg ctttatgcaa aacctaccac aatcgcttgc cgtggcttca ggagttgcat
56141 tttggataag tcaaaaagca agtgcttcta tattcgacaa gctctcgaat aaattagact cgaagattgc
56211 aattgctttg tctagcaaca tcggtttctat tatagacgca accatatgga tttcattag actgagtcct
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56351 tcttcagtc aattgcttcg agatatttga aaaagtagtc aggaaaattc ctgattatct tgcagtcaat
56421 tgcttcgaga tatttgaata agtagtcagg aaaattcctg attatttttt ttacaaaaac gcttgacttt
56491 attcattcat tattat

Table 29

Phage dp1 ORFs list

nb	Name	Frame	Position	Size (a.a.)	Key words
1	dp1ORF001	2	36698..40390	1230	Putative tail;
2	dp1ORF002	1	32386..35835	1149	Tail;
3	dp1ORF003	3	53538..55877	779	DNA polymerase I;
4	dp1ORF004	3	40401..42440	679	Minor structural;
5	dp1ORF005	1	23674..25434	586	
6	dp1ORF006	2	45296..46987	563	SWI/SNF Helicase;
7	dp1ORF007	3	22230..23621	463	Terminase;
8	dp1ORF008	1	49624..50961	445	DNAb Helicase;
9	dp1ORF009	2	13160..14404	414	
10	dp1ORF010	2	8699..9859	386	RecA;
11	dp1ORF011	3	28017..29096	359	Major head;
12	dp1ORF012	3	5346..6419	357	DNA pol. III beta;
13	dp1ORF013	3	10215..11240	341	DNA pol. III gamma and tau;
14	dp1ORF014	3	50961..51974	337	DNA primase;
15	dp1ORF015	1	3793..4728	311	
16	dp1ORF016	3	43413..44303	296	Amidase;
17	dp1ORF017	1	11242..12081	279	
18	dp1ORF018	3	35847..36686	279	
19	dp1ORF019	2	12161..12967	268	
20	dp1ORF020	1	1864..2658	264	exsD: Coenzyme PQQ;
21	dp1ORF021	2	2504..3295	263	GTP cyclohydrolase;
22	dp1ORF022	2	30896..31675	259	
23	dp1ORF023	2	6419..7195	258	
24	dp1ORF025	-1	18026..18778	250	
25	dp1ORF024	3	25992..26738	248	
26	dp1ORF026	2	21512..22252	246	
27	dp1ORF027	1	52762..53490	242	
28	dp1ORF028	3	44595..45299	234	
29	dp1ORF029	2	662..1348	228	exsB;
30	dp1ORF031	3	26943..27611	222	
31	dp1ORF030	-2	19423..20088	221	
32	dp1ORF032	1	52033..52647	204	
33	dp1ORF033	2	7670..8239	189	
34	dp1ORF035	-1	16859..17425	188	
35	dp1ORF036	1	48808..49362	184	DNAC replication;
36	dp1ORF037	1	55855..56388	177	
37	dp1ORF034	2	131..652	173	
38	dp1ORF038	3	1350..1871	173	exsC: 6-pyruvoyltetrahydropterin;
39	dp1ORF039	3	3306..3803	165	Citrulline biosynthesis;
40	dp1ORF040	1	7192..7683	163	
41	dp1ORF041	3	8208..8699	163	dUTPase;
42	dp1ORF042	1	48082..48561	159	
43	dp1ORF043	1	31699..32154	151	
44	dp1ORF044	-1	25211..25666	151	
45	dp1ORF045	2	25340..25777	145	
46	dp1ORF046	3	42774..43202	142	
47	dp1ORF047	1	47542..47961	139	
48	dp1ORF048	-3	16308..16709	133	
49	dp1ORF049	-3	43620..44018	132	
50	dp1ORF050	3	15081..15476	131	
51	dp1ORF051	2	29765..30154	129	
52	dp1ORF053	-3	49917..50300	127	
53	dp1ORF052	3	30516..30893	125	
54	dp1ORF054	2	14423..14800	125	
55	dp1ORF055	3	27627..28004	125	
56	dp1ORF056	-3	18780..19151	123	
57	dp1ORF057	1	9859..10218	119	
58	dp1ORF058	3	15633..15989	118	
59	dp1ORF059	1	30154..30507	117	
60	dp1ORF060	-2	37717..38070	117	
61	dp1ORF062	-3	44940..45284	114	
62	dp1ORF063	1	47200..47541	113	
63	dp1ORF064	2	29108..29449	113	

64	dp1ORF066	-3	28566..28898	110	
65	dp1ORF067	-1	44735..45061	108	
66	dp1ORF068	3	29451..29768	105	
67	dp1ORF069	-3	20094..20411	105	
68	dp1ORF061	-3	19161..19475	104	
69	dp1ORF070	1	15973..16284	103	
70	dp1ORF071	3	38904..39209	101	
71	dp1ORF072	-2	50749..51045	98	
72	dp1ORF073	3	14262..14555	97	
73	dp1ORF074	3	32298..32591	97	
74	dp1ORF075	-1	22154..22447	97	
75	dp1ORF076	-1	5435..5728	97	
76	dp1ORF077	1	14800..15084	94	
77	dp1ORF079	-3	35007..35288	93	
78	dp1ORF081	-3	55188..55466	92	
79	dp1ORF103	2	49352..49627	91	
80	dp1ORF080	1	42490..42759	89	
81	dp1ORF082	1	44728..44994	88	
82	dp1ORF083	-1	35720..35974	84	
83	dp1ORF065	-3	51246..51497	83	
84	dp1ORF085	-3	10602..10847	81	
85	dp1ORF087	-2	29794..30036	80	
86	dp1ORF088	3	5040..5279	79	
87	dp1ORF089	-2	12256..12495	79	
88	dp1ORF273	3	56256..56486	76	
89	dp1ORF078	-3	17280..17507	75	
90	dp1ORF090	1	27037..27261	74	
91	dp1ORF091	1	43189..43413	74	Holin;
92	dp1ORF092	3	46989..47213	74	
93	dp1ORF093	-2	45538..45756	72	
94	dp1ORF095	3	8877..9089	70	
95	dp1ORF096	-1	46469..46681	70	
96	dp1ORF097	-1	38888..39100	70	
97	dp1ORF098	1	43627..43836	69	
98	dp1ORF099	3	38298..38507	69	
99	dp1ORF100	1	1597..1803	68	
100	dp1ORF101	2	19220..19426	68	
101	dp1ORF094	1	8281..8484	67	
102	dp1ORF102	2	4034..4237	67	
103	dp1ORF104	-1	21224..21427	67	
104	dp1ORF105	-2	1828..2028	66	
105	dp1ORF106	-3	10329..10529	66	
106	dp1ORF108	-1	49250..49447	65	
107	dp1ORF109	-2	31435..31632	65	
108	dp1ORF110	1	16444..16638	64	
109	dp1ORF111	1	28657..28851	64	
110	dp1ORF113	-2	17521..17715	64	
111	dp1ORF084	1	15445..15636	63	
112	dp1ORF114	2	52952..53143	63	
113	dp1ORF115	-3	5151..5342	63	
114	dp1ORF116	-1	20474..20662	62	
115	dp1ORF117	-3	24492..24680	62	
116	dp1ORF118	2	15023..15208	61	
117	dp1ORF119	2	41054..41239	61	
118	dp1ORF120	1	28387..28569	60	
119	dp1ORF121	3	39222..39404	60	
120	dp1ORF122	-1	40220..40402	60	
121	dp1ORF123	-2	21145..21327	60	
122	dp1ORF124	-3	17712..17891	59	
123	dp1ORF125	-3	49740..49916	58	
124	dp1ORF126	-3	15960..16136	58	
125	dp1ORF127	-3	13335..13511	58	
126	dp1ORF128	1	4852..5025	57	
127	dp1ORF129	2	25133..25306	57	
128	dp1ORF130	-1	16619..16789	56	
129	dp1ORF131	1	43846..44013	55	
130	dp1ORF132	-1	15137..15304	55	
131	dp1ORF133	-2	7900..8061	53	
132	dp1ORF135	3	780..938	52	
133	dp1ORF136	-1	55094..55252	52	
134	dp1ORF137	-2	36988..37146	52	

135	dp1ORF138	-3	30504..30662	52	
136	dp1ORF139	-3	11934..12092	52	
137	dp1ORF140	3	20562..20717	51	
138	dp1ORF141	-1	42767..42922	51	
139	dp1ORF142	-3	31743..31898	51	
140	dp1ORF143	-3	7410..7565	51	
141	dp1ORF144	1	36517..36669	50	
142	dp1ORF145	1	42067..42219	50	
143	dp1ORF146	1	51484..51636	50	
144	dp1ORF147	1	55207..55359	50	
145	dp1ORF148	-1	28484..28636	50	
146	dp1ORF150	-3	15033..15185	50	
147	dp1ORF134	-2	349..498	49	
148	dp1ORF151	1	28027..28176	49	
149	dp1ORF152	1	42235..42384	49	
150	dp1ORF153	2	22307..22456	49	
151	dp1ORF086	2	52760..52906	48	
152	dp1ORF154	2	18446..18592	48	
153	dp1ORF155	3	13512..13658	48	
154	dp1ORF156	3	18777..18923	48	
155	dp1ORF157	-2	13135..13281	48	
156	dp1ORF158	-3	40581..40727	48	
157	dp1ORF159	-3	30225..30371	48	
158	dp1ORF149	-3	26331..26474	47	
159	dp1ORF160	2	41324..41467	47	
160	dp1ORF161	2	52175..52318	47	
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162	dp1ORF163	3	40224..40367	47	
163	dp1ORF164	-2	6553..6696	47	
164	dp1ORF165	-3	50361..50504	47	
165	dp1ORF166	-3	23376..23519	47	
166	dp1ORF167	3	1008..1148	46	
167	dp1ORF168	-2	54205..54345	46	
168	dp1ORF169	-2	45814..45954	46	
169	dp1ORF170	-2	27460..27600	46	
170	dp1ORF171	-3	47538..47678	46	
171	dp1ORF172	-1	10325..10462	45	
172	dp1ORF173	-2	32023..32160	45	
173	dp1ORF174	-2	29629..29766	45	
174	dp1ORF175	-2	15511..15648	45	
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176	dp1ORF177	-3	19800..19937	45	
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180	dp1ORF180	-1	41042..41176	44	
181	dp1ORF181	-1	12992..13126	44	
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183	dp1ORF183	-2	13762..13896	44	
184	dp1ORF184	-3	53196..53330	44	
185	dp1ORF185	1	22522..22653	43	
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187	dp1ORF187	2	34415..34546	43	
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189	dp1ORF189	2	42587..42718	43	
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193	dp1ORF193	-2	42325..42456	43	
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195	dp1ORF195	-3	42453..42584	43	
196	dp1ORF196	-3	11142..11273	43	
197	dp1ORF107	1	10750..10878	42	
198	dp1ORF197	2	7484..7612	42	
199	dp1ORF198	2	24119..24247	42	
200	dp1ORF199	-1	15614..15742	42	
201	dp1ORF200	-3	47715..47843	42	
202	dp1ORF201	1	38569..38694	41	
203	dp1ORF202	2	44483..44608	41	
204	dp1ORF203	-3	22656..22781	41	
205	dp1ORF204	1	1471..1593	40	

206	dp1ORF205	1	8524..8646	40	
207	dp1ORF206	1	19855..19977	40	
208	dp1ORF207	1	27502..27624	40	
209	dp1ORF208	2	47279..47401	40	
210	dp1ORF209	3	29784..29906	40	
211	dp1ORF210	-1	52955..53077	40	
212	dp1ORF211	-1	20837..20959	40	
213	dp1ORF212	-2	52861..52983	40	
214	dp1ORF213	-2	30169..30291	40	
215	dp1ORF214	-2	24151..24273	40	
216	dp1ORF215	-3	35700..35822	40	
217	dp1ORF216	-3	32727..32849	40	
218	dp1ORF217	1	23443..23562	39	
219	dp1ORF218	3	22029..22148	39	
220	dp1ORF219	-1	51269..51388	39	
221	dp1ORF220	-1	6215..6334	39	
222	dp1ORF221	1	43507..43623	38	
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226	dp1ORF225	-2	32875..32991	38	
227	dp1ORF226	-2	25075..25191	38	
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229	dp1ORF228	1	10450..10563	37	
230	dp1ORF229	1	27634..27747	37	
231	dp1ORF230	2	50723..50836	37	
232	dp1ORF231	-2	30958..31071	37	
233	dp1ORF232	-2	29272..29385	37	
234	dp1ORF233	-3	52779..52892	37	
235	dp1ORF234	1	36253..36363	36	
236	dp1ORF235	2	32768..32878	36	
237	dp1ORF236	-1	37418..37528	36	
238	dp1ORF237	-1	1568..1678	36	
239	dp1ORF238	-3	1191..1301	36	
240	dp1ORF239	1	26521..26628	35	
241	dp1ORF240	1	41893..42000	35	
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248	dp1ORF247	1	29641..29745	34	
249	dp1ORF248	1	53560..53664	34	
250	dp1ORF249	2	2012..2116	34	
251	dp1ORF250	2	23837..23941	34	
252	dp1ORF251	-1	39101..39205	34	
253	dp1ORF252	-2	54667..54771	34	
254	dp1ORF253	-3	56151..56255	34	
255	dp1ORF254	-3	48375..48479	34	
256	dp1ORF255	-3	9468..9572	34	
257	dp1ORF256	1	15289..15390	33	
258	dp1ORF257	1	28216..28317	33	
259	dp1ORF258	1	44023..44124	33	
260	dp1ORF259	2	4298..4399	33	
261	dp1ORF260	2	24746..24847	33	
262	dp1ORF261	3	288..389	33	
263	dp1ORF262	3	9408..9509	33	
264	dp1ORF263	-1	26951..27052	33	
265	dp1ORF264	-1	6038..6139	33	
266	dp1ORF265	-1	4700..4801	33	
267	dp1ORF266	-2	50119..50220	33	
268	dp1ORF267	-2	47266..47367	33	
269	dp1ORF268	-2	12520..12621	33	
270	dp1ORF269	-3	53733..53834	33	
271	dp1ORF270	-3	50691..50792	33	
272	dp1ORF271	-3	19638..19739	33	
273	dp1ORF272	-3	1455..1556	33	

Table 30

Predicted Dp-1 amino acid sequences

dp1ORF001

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36866 gaaacctcatctatctatcaacacttaaggttgaacacattatccagatggaggaagatgggttcgaattaaatagctcag
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36950 gacgtagaagatgtcaagggttaccagtttacctgctacgcattatggatgaactagcagaaggcttgcctaggaattg
85 D V E D V K G L T K F T C Y A L W Y E L A E G L P R K L
37034 aaacacgttgcctctctgtaggcgctgtcgcgctagatattatcaaagacgcaggatgaatgggttcgactagttgtcctcct
113 K H V A S S V G A V A L D I I K D A G E W V R L V C P P
37118 gacggtgctaaacaaagttcgaagcataacagccgcagaaaattcaatgctttggcattcttcgatatcttgcgaagcaatac
141 D G A N K Q V R S I T A A E N S M L W H L R Y L A K Q Y
37202 aatttagaattgacatttgggtatgaagaattatcaagcaagaggttagaattgttcaaacgttgcatttcttcagccttat
169 N L E L T F G Y E E I I K Q E V R I V Q T V V F L Q P Y
37286 gtcgagttctaaagtagactttcctcttggtagtgaagagaatttgaatatgtcactaggcaggaagattctcgaacacctgtgt
197 V E S K V D F P L V V E E N L K Y V T R Q E D S R N L C
37370 acggcttacaagttgacaggtaaaaaggaagagcagtcgaagacgttcaacgtttgcttctatcaacaatggaagtgaaatat
225 Y N K V P D L H H T Q L I V D D H Y D V I E W R K I S A
37454 ctcatgatgtttcgtggttactacacgccacatgaagcctcgatatattgctaattctaaaagcgaacacattttagaatt
253 L I D V S W F T T R H M K P R Y I A K S K S D E H F R I
37538 aaagaaaatttgatgagtgctgcgcgtgcttcttgcacatctacagtcgccactaattggatatgaggttccagcggctcctt
281 K E N L M S A A R A Y L D I Y S R P L I G Y E A S A V L
37622 tataacaaggttcctgacttgcacatactcaactaattgtcgacgaccattatgatgttatcgagtgaggaaagatatctgct
309 Y N K V P D L H H T Q L I V D D H Y D V I E W R K I S A
37706 cgaaaaattgactacgacgacctttcaactctactatcattttcgaagacccctcgaaaagacttgatggacttgcataatgag
337 R K I D Y D D L S N S T I I F Q D P R K D L M D L L N E
37790 gacggcgagggagtcctttcaggggaactgttaattgactcccaagttgttatttagatagcagatgacatttttagggactaat
365 D G E G V L S G E T V N E S Q V V I R Y A D D I L G T N
37874 tttaatgcagaatctgggaatacattgggtgccttaactaataaagaaacgcagcgaattagttcctgacgactttacatgg
393 F N A E S G K Y I G V L N T N K K P S E L V P D F T W
37958 attcgcactagaaggtcctaaaggtgacgcaggtttaccgggagctcctggcggtgatggagtcgacggtgtacctggaaagagc
421 I R L E G P K G D A G L P G A P G R D G V D G V P G K S
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449 G V G I A D T A I T Y A V S V S G T Q E P E N G W S E Q
38126 gttcctgaactcataaaaggtcgattcctgtggactaaaacattttggagatatactgacggtcactgaactggatgactcct
477 V P E L I K G R F L W T K T F W R Y T D G S H E T G Y S
38210 gttgcctatatagggcaagacggaaattccggaaagacggaatcgacggttaaggacggagtaggtatagccgcaactgaagtc
505 V A Y I G Q D G N S G K D G I A G K D G V G I A A T E V
38294 atgtatgcaagttcgccatctgctactgaagctccagctgggtggatgggtctacgcaagttcctaccgtcccaggtgggtcagtat
533 M Y A S S P S A T E A P A G G W S T Q V P T V P G G Q Y
38378 ttatggactcgaacaagatggcgctacactgacaaactgtgaaattggatattcagtttcaagaatgggcgagcaggttcct
561 L W T R T R W R Y T D Q T D E I G Y S V S R M G E Q G P
38462 aaaggtagcgcaggtcgtgacggtattgcaggaagaaacggaataggggttaagtcacactcagtttcttggaaattagtcctc
589 K D A G R D G I A G K N G I G L K S T S V S Y G I S P
38546 actgattctgcgattcctggagtatgggcttcacaagttccttcttaatacaagggtcaatatctttggactcgaactatttgg
617 T D S A I P G V W A S Q V P S L I K G Q Y L W T R T I W
38630 acctataccgattcaactaccgaaacgggctatcaaaaaacctacattccaaaagacgggaatgacggttaaaaatggaattgct
645 T Y T D S T T E T G Y Q K T Y I P K D G N D G K N G I A
38714 ggtaaggatgggttaggaattaagtctacgaccattacctacgcaggctcaacctcaggaacagttgcgctacttcaaattgg
673 G K D G V G I K S T T I T Y A G S T S G T V A P T S N W
38798 acttctgctattccaaatgttcaaccgggattcctcttggagcgaacactgtttggaactatactgatgacactgacgaacaa
701 T S A I P N V Q P G F F L W T K T V W N Y T D D T S E T
38882 gggttactcagtttccaaagataggtgaaacaggtcctagaggagttcaaggcttccaaaggtcctcaagggttccaaaggttcc
729 G Y S V S K I G E T G P R G V Q G L Q G P Q G L Q G I P
38966 ggacctgcaggagctgacggacgttcgcaatatactcacctcgcttctcctaattagtcctcaaacgggtgaggttagtcatact
757 G P A G A D G R S Q Y T H L A F S N S P N G E G F S H T
39050 gacagcggacgagcagtcagtcaggtcagatcaagatttcaatcccgctccattcaaaagaccctgcagcctatatactggacgaaa
785 D S G R A Y V G Q Y Q D F N P V H S K D P A A Y T W T K
39134 tggaaaggggaatgacggagctcaagggtatcccggaagccaggcgacagcggtaagactaattatttccatatagcttacgct
813 W K G N D G A Q G I P G K P G A D G K T N Y F H I A Y A
39218 tcaagtgcagacggatcacgtgagttcagtttgggaagataatacaacaatatatgggttattactccgattatgagcaafca
841 S S A D G S R E F S L E D N N Q Q Y M G Y Y S D Y E Q A
39302 gatagcagggatcgaaactaagatcgatgggttgaccgcttgcaaatgttcaagtgagggtcgaaacgagttccttaattct
869 D S R D R T K Y R W F D R L A N V Q V G G R N E F L N S
39386 ttatttgaatttgggttaaaacctcgctattctagttacaatctaattggacggacaagatcaaacgcaaggacagatatctgct
897 L F E F G L K P R Y S S Y N L M D G Q D Q T Q G Q I S A
39470 actattgacgacgctcaacgggttcaaggttgcactaactcttgcagacttgactcaacatggaaacggtaaacccgagaacccaaaa
925 T I D E R Q R F K G A N S L R L D S T W N G K P Q N Q K

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39554 ctgacccctttcttaggagagatagcgattaggtactccaaccgagtggtctaatcttagaaggctcgatcttctgggct
953 L T F S L G G D T R L G T P T E W S N L E G R I S F W A
39638 aaggcccttaggaacggagtgagcttagctgacggcggttatcgtagtaacgtatttaccgcaaccttaaccgatcaatgg
981 K A S R N G V S L A A R P G Y R S N V F T A T L T D Q W
39722 aagttctacgattttaaattctttgacaaaagtttaattgacggctgaagcaattttccatgtattcactcaaatgtgt
1009 K F Y D F K F F D K V N S N C T A E A I F H V F T Q S C
39806 tcagttgtggctcaatcatattaaaatcgaaacttggttaatatctctactccttttagtgaagcagaggaaagaccttaaatatcga
1037 S V W L N H I K I E L G N I S T P F S E A E E D L K Y R
39890 attgactcaaaagccgatcaaaagctaactaaccaacagttgacggcactcacggaagggctcaactacatgacgcagaactg
1065 I D S K A D Q K L T N Q Q L T A L T E K A Q L H D A E L
39974 aaagctaaggctacaatggagcagtttaagtaacttagaaaaggcttatgaaggtagaatgaaagctaataagaagctatcaaa
1093 K A K A T M E Q L S N L E K A Y E G R M K A N E E A I K
40058 aaatcggaagccgacctaattcttagcggaagtcgaattgaagctactatccaagaacttgccgggtacgggaactgaagaag
1121 K S E A D L I L A A S R I E A T I Q E L G K L R E L K K
40142 ttctgacgacgttacatgagctcttctaatgaaggtctaattatcggtgaagaacgacggtagctctaccattaaggtatcaagt
1149 F V D S Y M S S S N E G L I I G K N D G S S T I K V S S
40226 gaccgaatttctatgttctcgcaggggaatgaagttatgtaccttacgcaagggtcattcacatcgataacgggacattcttacc
1177 D R I S M F S A G N E V M Y L T Q G F I H I D N G I F T
40310 caatccattcaagtcggcgttagaagcgaacaatactcggttaataccagacatgaacgtgattcggtatgtaggataa
40390
1205 Q S I Q V G R F R T E Q Y S F N P D M N V I R Y V G *
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29 R A L A L E S S K S F Q I G S A L T G L G K G L T T A V
32554 acccttctcttatgggatttgacggcctctattaaagtaggggaatgaattccaagctcaaatgtccgtgttcaagctatt
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32722 gcggctcaaggtatggaatctagcttcagcgggttccaggttaaatgaaatcatggacgctatgccaggggtacttgacctg
113 A Q G M E N L A S A G F Q V N E I M D A M P G V L D L
32806 gctccgtatctggagagatgtggcgcgagctccagggccatggctagttcacttcgagcctttggattagaggcaaacag
141 A V S G G D V A A S S E A M A S S L R A F G L E A N Q
32890 gcgggtcacgtggctgacgtatttgctcgagcagcagctgatacgaacgcagaaactagcgacatggcagaggcgatgaaatc
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197 V A P V A H S M G L S L E E T A A S I G I M A D A G I K
33058 ggctcgcaagccggaacacagcttagaggcgctctctcgcgtattgccaaacctacgaagcgatgggtcaaatcaatgcaggaa
225 G S Q A G T T L R G A L S R I A K P T K A M V K S M Q E
33142 ttaggagtttctgctacgacgcgaacggaacatgattccactaagagaacaaatcgctcaactgaaacagctactgcagga
253 L G V S F Y D A N G N M I P L R E Q I A Q L K T A T A G
33226 ctaacacaaggaagcaaaaactcgtaccttggctacgttatggcctcaaaactcggtgctcaggtatgcttcgactatgacgca
281 L T Q E E R N R H L V T L Y G Q N S L S G M L A L L D A
33310 ggtcctgagaaattggataagatgaccaatgctctcgtgaactcgacggagctgctaaggaaatggcagaaactgcaggac
309 P E K L D K M T N A L V N S D G A A K E M A E T M Q
33394 aaccttgctagtaaaatcgagcaaatgggaggagctttcgagctgctgctattattgttcaacaaatccttgagcctgcact
337 N L A S K I E Q M G G A F E S V A I I V Q Q I L E P A L
33478 gctaaaatcgtagggagcaatcacaagcttctcgaagctatcgtaaatatgtcacctatcggtcaaaagatgggtgctcatattc
365 A K I V G A I T K V L E A F V N M S P I G Q K M V V I F
33562 gcaggaatgggtgcagcccttgaccactgcttcaattgcaggaatgggtgatgacaactattgtcaagtaagaattgctatt
393 A G M V A A L G P L L L I A G M V M T T I V K L R I A I
33646 cagtttttaggtccagcatttatgggaacgatgggaacattgcaggagttatagcaatattctatgctcgtgctggcgtgcttc
421 Q F L G P A F M G T M G T I A G V I A I F Y A L V A V F
33730 atgatgcctacacaaaatcgagagatttagaaaactttatcaacagctcttgccctgctattaaagctgggtttggaggagcg
449 M I A Y T K S E R F R N F I N S L A P A I K A G F G G A
33814 ttggaatggctacttccacgactgaaagagtttaggagaatgggttacagaaggcaggcgagaaggcgaaagagctcggtcagct
477 L E W L L P R L K E L G E W L Q K A G E K A K E F G Q S
33898 gtagggtctaaagtgtcaaaactgctcgaacagtttggaataagtatcggtcaggcaggaggtcgattggtcagttcattgga
505 V G S K V S K L L E Q F G I S I G Q A G G S I G Q P I G
33982 aatgttctcgaaaggctaggaggcgctatttggaagtaggaggagtcatttcaattgctgtttcacttgtaacaaaattcggt
533 N V L E R L G G A F G K V G G V I S I A V S L V T K F G
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701 L V Q A L P T L I I A T T A C G T Q I L S A L I N G L V Q A L P
34570 gcaattattcaagcagctgttcaaatatcagtcgctgttcaagcactaattgaaaacttgctctatgataatcgaagcagc
729 A I I Q A A V Q I I M S L V Q A L I E N L P M I I E A A

365

34654 atgcagattataatgggtctagtcacgcactgattgaaaataggacattcttagaagcagggttcaaattctaatggct
757 M Q I I M G L V N A L I E N I G P I L E A G I Q I L M A
34738 ttaatcgagggtacttattcaagtgtcttctgaactaattacagcagcgattcaaattcattacttcaactattagaagcaattcttg
785 L I E G L I Q V L P E L I T A A I Q I I T S L L E A I L
34822 tcgaaccttctcaacttctagaagcggagttaaattgcttttctcacttcttcaagggttgctaaatagcttctcactaacta
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34906 attgcaggggttgcgaatcatgatggcacttcttaagcagttatcgacttcgctcctaaacttcttcaagcaggtgttcaa
841 I A G A L Q I M M A L L K A V I D F V P K L L Q A G V Q
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869 L L K A L I Q G I A S L L G S L L S T A G N M L S S L V
35074 agcaagattgttagcttctgtgggacagatgggttcaggagggtgcgaacctgattcgaaacttcattagtggtattgggtcaatg
897 S K I A S F V G Q M V S G G A N L I R N F I S G I G S M
35158 attggttcagctgtctctaaattggcagcatgggaacttcaattgtttctaaagggttactggattcgctggcaaaatggtaagc
925 I G S A V S K I G S M G T S I V S K V T G F A G Q M V S
35242 gcagggttcaaccttctcgaggatttattcaatgggtatcagtttccatggtaagttctgctggtaagtcggtcgtcgaatggct
953 A G V N L V R G F I N G I S S M V S S A V S A A A N M A
35326 agcagtgcatataatgccgttaagggttcttaggtattcacttctcctcagctgtcatggagcagatgggtattctatcagggt
981 S S A L N A V K G F L G I H S P S R V M E I Y T G
35410 caagggttcgtaaatggtattggttaacatgattcgaactacacgtgacaaaggctaaagaaaggctgaaactgttactgaagct
1009 Q G F V N G I G N M I R T T R D K A K E M A E T V T E A
35494 ctcagcgacgtgaagatggatattcaagaaatggagttatagaaaagggttaaatcagtttaccgaaagatgggtgaccaactt
1037 L S D V K M D I Q E N G V I E K V K S V Y E K M A D Q L
35578 cctgaaactcttcagctcctgatttgcgaagatgttcgtaaaagcagcgggttcgctcagtgaggttcttcaatacaggaagt
1065 P E T L P A P D F E D V R K A A G S P R V D S L F N T G S
35662 gacaaacctaaccaactcagtcacacttcaaaacatcaaggcgagcaaacggttctgcaacttggaacaactcgttagttcga
1093 D N P N Q P Q S Q S K N N Q G E Q T V V N I G T I V V R
35746 aacaatgacgaggttgacaaactgtcgagaggattgtataatagaagtaagaaactctatcagggtttggtaacattgtaaca
1121 N N D D V D K L S R G L Y N R S K E T L S G F G N I V T
35830 ccgtaa 35835
1149 P *

dp1ORF003

53538 atggcacaataaggactcttgggtgcaagcctcggtctagcaagaagaacgatgctcagttacttctcaacggaaaaacagg
1 M A Q K G L F G A K P R S S K K N D A Q L L A Q R K N R
53622 aagcctgcagttgaggttacttacttcaacttcaaggaaacgctctaaaggacgagttgcttagagctcgtacttctcaactaggatt
29 K P A V E V T Y I S G N A L K D A V A R A R T L S T R I
53706 cttggacacgttcttgatagacttgagttaatcactgaggaagcaaaactcgagcagttatgtagacaaaatgattgaagacgga
57 L G H V L D R L E L I T E E A K L E Q Y V D K M I E D G
53790 atagggttctattgacgtgaaacttgaggtcagttcactcagatgagctggcagaggtctgcttgactcactcaggtcactcag
85 I G S I D V E T D G L D T I H D E L A G V C L Y S P S Q
53874 aaaggaattctatgctcctgtcaatcatggttagcaatagacgaagatgcgaattagaatcaaatcttctcctgagttcatgaag
113 K G I Y A P V N H V S N M T K M R I K N Q I S P F M K
53958 aaaaatgcttcaacggttggatgattcaggaattcctgtcactatcataattcgaaatttgacatgaatcgatttattggcga
141 K M L Q R I V D S G I P V I Y H N S K F D M K S I Y W R
54042 ctcggcgtcaaaatgaatgagccagcgtgggtatcatatttagccgcaatgcttttaaatgaaaacgagtcacagcttgaaa
169 L G V K M N E P A W D T Y L A A M L L N E N E S H S L K
54126 agtcttctacttaaatgatttaggaacgaagaacgcagaggttgcaaaatttaattgacttatttaagggaattcctttagt
197 S L H S K Y V R N E E N A E V A K P N D L F K G I P F S
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225 L I P P D V A Y M Y A A C C Y D P L Q T F E L Y E Y D
54294 ttgactccaggaactgaacaatgtgaagaatataaactggaaaaagttctcatgggttcttcataatattgagatgcctctaatt
253 L T P G T E Q C E E Y N L E K V S W V L H N I E M P L I
54378 aaagttcttctcgacatggaagtctacggttcgacttagaccaagataagctggcagaattagagaacagtttactgccaat
281 K V L F D M E V Y G V D L D Q D K L A E I R E Q F T A N
54462 atgaacgaggtgagcaagagtttcaacagcttctgcagcaatggcagcctgaaattgaagaacttcgacaaactaatttccag
309 M N E A E Q E F Q Q L V S E W Q P E I E E L R Q T N F Q
54546 agctatcaaaaactcgaaatggatgcaagaggtcgagtgacggttaagcatttccagtcctactcaattagcaattctgtttat
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54630 gatattcatgggttgaaggtcctgaaagggtataaaccttagaggaacagggcgaaggtattgtcgagcattttgataacgatattc
365 D I M G L K S P E R D K P R G T G E S I V E H F D N D I
54714 tcaaaagcacttttgaatatagaaaatagcaaaatagtttgcacctatatacaacttgaccaacaccttgcaagcctgac
393 S K A L L K Y R K Y A K L V S T Y T T L D Q H L A K P D
54798 aatcgaattcacactacattcaaacagtcagggagctaaagacagggcggtatgtcaagtgagaatcctaacttacagaatattcct
421 N R I H T T F K Q Y G A K T G R M S S E N P N L Q N I P
54882 tctcgcggtgaggggtcagtagttcgacaaacttctgcagccagtgagggttacttattggttagtgactacttcaacaa
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55134 agaaattctgtcaagtcggttcttttaggttcttatgtacggcgggggttaactcaatcgctgagcagatgaattgattctgtc
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55218 aaagaagcgaataaggttattgaagatttcttccagagcttccctaaagtgaggcagactatcatattcgttcaacagcagcg
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55302 caggacttgggtatgttcaaacagctaccggtcgaagaagaagggttctctgatattgattcttccgtaacagaggttcaggtat
589 Q D L G Y V Q T A T G R R R R L P D M S L P E Y E F E Y
55386 atcgacgctagcaagaacgaagatttcgacccctttaaactttgacgcagaccaacagatggacgatactgttctgacatatt

366

617 I D A S K N E D F D P F N F D A D Q Q M D D T V P E H I
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 645 I E K Y W A Q L D R A W G F K K K Q E I K D Q A K A E G
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 253 L Q I M S N I Q V N P N N A S G A Y G S T I Q A P H A E
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 393 G T F T T I S L M T N S S A N L A G N Y G P D K S Y I V
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 673 D N V S F R I *
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 113 R I L F D S I L R N C K F W S K S T N A L V D A T V G K
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 141 R V L M T V V A N A A Q Q I D V Q F Y S M P Q F T Y T V
 24178 gaccctagaaaaccccttcagcttcttctgttgacattgtttatcaggacgagcgtacaaaaggaatgagcactgaaaaacaa

367

169 D P R N P S S L L S V D I V Y Q D E R T K G M S T E K Q
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197 L W H H Y R Y E M K A G T S Q S G I A T A L E D I E E Q
24346 tgttggctcacttatgccttaacggatggagagtcgaacaaatctatatgacagaaagtgccaaactactatcaaggagaca
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365 G T G G K Q A Q V T S I S G N P N F L P A E I S E
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393 G A K K A M Y E L M D Q P M P E K V Q E A P S G I A M Q
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421 F L F Y D L I S R C D G K W I E W D D A I Q W L I Q M L
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449 E I L A T V N V D L G N I P Q D I Q S T Y Q T L T M
25102 actatcgaaacaccactatccaattcctagcgagaaacttctctgaagcaacttgcgctcactgaagttcaactaatgtacgc
477 T I E H H Y P I P S D E L S A K Q L A L T E V Q T N V R
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25270 cttgacgaatctcagctggagcattgcctgtattagcaaacgaatataacgaacaagaggagcctcaagatgaacagagtgaa
533 L D E I S A G A L P V L A N E L N E Q E E P Q D E T S E
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25434
561 E D E V D D K E K E Q T E Q P T E E G V D P D V Q G *
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57 F N N V I D A L D E W E L H I F G E L D K D V Q D Y I D
45548 tctcgaaacgaatagcttcttcaagcaatgagcagtttgcgttcaagactactcattcgcgacccaggttgaatgttctgaa
85 S R N R I A S S S N E Q F S F K T T P F A H Q V E C F E

45632 tacgcacaagagcatccatgttctcttttaggcgagtagcaaggttttagggaaaactaaacaggcaattgatattgcagttagc
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45716 aggaaggcaagtttcaaacattgtttaatcgatcttgcatatcagggtcctaaatggaattgggcaaaagaagtaggtattcat
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169 S N E S A H I L G S R V T K D G K L V I D G V S K R A E
45884 gacttgccttgggtggccacgcaatcttcttactcaacattgaaactcttcgagtagctgttgcattaaataacttaaat
197 D L L G G H D E F F L I T T N I E T L R D A V F I K Y L N
45968 gaactgacaaaaagcgagaaattggaatgggtatttgacgagattcacaagtgaagaaccttcaagtaagcaaggggct
225 E L T K S G E I G M V I I D E I H K C K N P S S K G A
46052 tcaattcaaaagctccaaagtattacaagatgggacttacaggaaactcctctaataaatacccaatcgatgtattcaatgtt
253 S I Q K L Q S Y Y K M G L T G T P L M N N P I D V F N V
46136 atgaagtggttagggcggaacatcacactgactcagttcaagagcgatactgtatcgctgaccagttcaatcaaatcact
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46220 ggatattcgaaatctagctgaacttcgagcgttgtaacgactacatgcttagaagaacgaaggaaggttttagacctgcct
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46304 gaaaagattcgagtcacagagtagtgcgacatgaactcgaacagtcacaaatctataaggaagttttagctaaacttgcctcaa
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46388 gaaatagataaagtcgaagctcatgcctaacctctagccgaaacgattcgacttcgacaagcgactggaatccttcgatttta
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46976 ctgcttaaatag 46987
561 L L K *
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368

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169 T G S K M W F S C N P A N P N H Y F K K N W I D K Q V E
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197 K R I L Y L H F T M D D N P S L T D S I K R R Y E K M Y
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337 I R G K Q I E Y I I L D P S A S A M I V E L Q K H P Y I
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365 A R K N I P I I P A R N D V T L G I S F H A E L A E N
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393 R F T L D P S N T H D I D E Y Y A Y S W D S K A S Q T G
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169 G E D L I V I M A R P G Q G K S W T I D K M L A T A W K
50212 aacgggcatgatgtccttctatatagcggggaatgagtgaaatgcaagttggtgctcgtatagatactatttcttcgaatgtt
197 N G H D V L L Y S G E M S E M Q V G A R I D T I L S N V
50296 agcatcaattcaattacaaagggatttggaaagaccatcagttcgaaaaatagaggaccatattcaagcaatgactgaggct
225 S I N S I T K G I W N D H Q F E K Y E D H I Q A M T E A
50380 gaaaattcccttgtggtagtcacgccccttatgattggaggaaagaaccttacccttgcatttttagatagcatgatctaaa
253 E N S L V V V T P F M I G G K N L T P A I L D S M I S K
50464 tatagaccatctggtgggtgattgaccagctttcactcatgacgagcttatccaagcaggagcagaagcgaatccagtac
281 Y R P S V V G I D Q L S L M S E S Y P S R E Q K R I Q Y
50548 gccaacatcaccatggacctatataagatttctgctaaatattggaattcctattgtgcttaattgccaagcagggcggttcggct
309 A N I T M D L Y K I S A K Y G I P I V L N V Q A G R S A
50632 aaaactgaagcgctgaaagtattggaactagaacatatagcagaagtgatggagtaggtcaaaatgctagcagagttatcgct
337 K T E G A E S M E L E H I A E S D G V G Q N A S R V I A
50716 atgaagcgtgacgaaaaatccggcacttgaactatctgctgtaaaaacccgatatggcgaagacgaaaaatcatcgaatat
365 M K R D E K S G I L E L S V V K N R Y G E D R K I I E Y
50800 atgtgggacgttgaactggaacctatactcttataggattcaagaggaagggcgaagaaggaactgaaaaagggcgaagctct
393 M W D V E T G T Y T L I G F K E E G E E G T E K G E S S
50884 ccattgaaagcaaaagcctctaggtcgactgctcgtcttcgaagtaaggttacaaggggaagaggttgaagcattttga 50961
421 P L K A K A S R S T A R L R S K V T R E G V E A F *

dp1ORF009
13160 atgacagactttaaaaaacgcttcaagaaaagcagtaacagaaaacatcaatcgtagcgggtatcgagaaccttatggattggctc
1 M T D F K K R F K A V T E T I N R D G I E N L M D -W L
13244 gaaaatgataccaattttcttctcgaagcagcactcgataccatggaagctatgaaagtggaacttgcgagcactcatta
29 E N D T N F F S S P A S T R Y H G S Y E G G L V E H S L
13328 aacgtgttcaatcaactactttcgaatggataccatggtaggcaaggttgggaagacatttaccatggaaacagttgca
57 N V F N Q L L F E M D T M V V G K G W E D I Y P M E T V A
13412 atcgtagcactatttcacgaccttgcgaaggttgggtcagttcgtgaaactgaaaaatggcgcaagaacagcgaggtgaatgg
85 I V A L P H D L C K V G Q Y R E T E K W R K N S D G E W
13496 gaaagctatttagcatatgaatacgcacctgagcaacttacaatgggacatgggtgcaaaatctaatttcttctcaacgttctc

8699	atgaaattggaacggtgatgaaggactggaataaggattcgaaaagctcttctagcagttcaaggacttgaacgtgaagcgctt
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8783	ccaagaatccccctttctcgccctcttatgaattatcaaacctacgcggcctccctcgaaaaagggtagttgaaattcttcgggt
29	P R I P F S A A P S M N Y Q T Y G A G L P R K R V V E P F G
8867	cctgagtcgaagtggaagaaactactcagctctcgacattgtcgaagaatcgcgaaattggattttgagcaggaatgggaacagaa
57	P E S S G K K T T S A L D I V K N A Q M V F E Q E W E Q K
8951	actgaagaactcaagggaaaagctggaaaatcgcgctgcatacgaagctagcaagactgctgcaaggaacttgaatactgcaactc
85	T E E L K E K L E N A R A S K A S K T A V K E L E M Q L C
9035	gatagctcttcaagagcctcttaagattgtatatcttgaccttgagaatacattagacactgagtggtggctctaaaaagattggagtc
113	D S L Q E E P L K I V Y L D L E N T L D T E W A K K I G V
9119	gagtgctgacaattttggatagtttgcgcttgaattgaacagcgctcgcaagaataacttcaatatgttttagacattttcgaaaca
141	D V D N I W I V R P E M N S A E E I L Q Y V L D I F E T
9203	ggtagaagttggcctagtagttctagattctcctcgcttcatagcttcgagctctcaaaaccttattgatggaagattgactctcaaaagcc
169	G E V G L V L V L D S L P Y M V S Q N L I D E E L T K K A
9287	tatgcaggaatctcagcgcccttctgactgaatttagctcgaaaggttactcctctcttactcgctacaaatgcaatattctctaggc
197	Y A G I S A P L T E F S R K V T P L L T R Y N A I F L G
9371	atcaatacaattcgagaagatatgataagtcagtaacattgcctattcaccatccgaagcggaagattggaagcatgctctgtgca
225	I N Q I R A E D M N S Q Y N A Y S T P G G K M W K H A C A
9455	gttcgacttaaaattgaaaaaggtdgactaccttgacgaaaacgggtgcattcattgaccccgctactgctcgaaacccctcaggggaat
253	V R L K F R K G D Y L D E N A G A S L T R T A R N T P A G N
9539	gtagttagagtcattcgtcgagagacccaagcatttaagcgcgacagaaaattagtttctctatcgcgttctctatcatgatgga
281	V V E S F V E K T K A F K P D R K L V S Y T L S Y H D G
9623	attcaaatgtaaaatgaccttgtagatgctcgctgtcgaatttggagtcattcaaaaggcaggggcatgggttcagtatcgtcgac
309	I Q I E N D L V D V A V E F G V I Q K A G A G A W F S I V D
9707	ctgaaactggagaaaattgacagatgaagacgaagaaccattgaggttccaagggcaaggcaaatctagttcgacgcttcaag
337	L E T G E E I M T D E D E E P L K F Q G G K A N L V R R F K
9791	gagtagtactacttctcgacatgggtgatgactcggttccacgaatttatcactcgagagaagaggtcaa 9859
365	E D D Y L F T D M V M T A V H E I I T R E E G *

28017	atgaatatttatgattatatcaacgcaggggagattgctagctacattcaagcacttcttccaacgctcttcaaatccttgga
1	M N I Y D Y I N A G E I A S Y I Q A L P S N A L Q Y Y L G
28101	ccaactctttdtccctaatgctcaacaacaggagacacatttctggctcaagggtgcaaatcaatttgcgtaactatccag
29	P T L F P P N A Q Q T G T D I S W L K G A N N L P V T I Q
28185	ccatctactacgcgcgaagcaagctctctgtgacagctgtggatttagcaacaacagctactgagatggcattcttccgtgag
57	P S N Y D A K A S L R E R A G F S K Q A T E M A F F R E
28269	tctatgcgacttgggtgaaaaagaccgtcaaaacttgcaaatgctattgaaccaaagttcagctcttgcaccaaccacttatcact
85	S M R L G E K A C D R Q N L Q M L L N Q S S A L A Q P L I T
28353	caactctataatgatactaagaaccttgtagacggtgttgaaagcgcaagcaatacatcgctatgcaattgctctcaatccggt
113	Q L Y N D T A C T K N L V D G C G V T T G A A Q A E Y M R M Q L L Q Y G
28437	aaattcactgtccaatcaactaacacgcagggtcctaactctacgactcaacactggatgctgaagcaacaacttgcagctcact
141	K F T V K S T N S E A Q Y T Y D Y N M D A K Q Q Y A V T
28521	aagaatggactaacccagctgaaagtgaacctatcgctgacattttagcagcaatggatgacatcgaaaatcgtaacaggtgtt
169	K K W T N P A E S D P I A D I L A A M D D I E N R T G V
28605	cgccctactcgaattggtcttgaaccgaaacacttataaccaatgactaagatgactctatcaagaagaactcttgcgaattggt
197	R P T R M V L N R N T Y N N Q M T K G A S D S I K K A L A I G
28689	gttcaaggtctctgggaanaactcttctgttcttgcaagtgacgtgagaaattcactcgctgaataaacaggctcttcaactcga
225	V Q G S W E N F L L L A S D A E K F I A E K T G L Q I A
28773	gtctactctaagaaaattgctcagttcgctgacgtgacaaaacttctgacgttggtaacattcgtcagttcaacttgattgac
253	V Y S K K I A Q F A D A D K L P D V G N I R Q F N L I D
28857	gacggtgaagtggtattgcttccactgacgcagcttggctcacacttggtagctactactcagaagcattcgacttggcttca
281	D G G K V V L L P P D A V G H T V Y G T T P E A F D L A S
28941	ggcggaacagcgtcgaagttcaagttcttctcaggcggaacctaccggttacaacttacttggaaaaaacctctgtgcaacattgca
309	G G T D A Q V V Q V L S G G P T V T T Y L E K H P V N I A

29025 acagttgtatcagctgttatgattccatcattcgaaggaattgactatgtaggagttctcacactaattag 29096
337 T V V S A V M I P S F E G I D Y V G V L T T N *
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5346 atgagtattaaagttcaaaacgaagaactttcaaaaattgtttctcagctcaataagttgaagcctagcaagttgctagaatc
1 M S I K F K T E E L S K I V S Q L N K L K P S K L L E I
5430 acaaaattattggcatatttttggtagcggcgaatgcgtcatgtttacagcgtatgatggctcaaaacttcctcgatgcattatc
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5514 gacagcgtatgtgaaattgacgtgattgtgaaagcagagcagtttggaaaactgtgaaaaagaccagcgccgcaacttcacac
57 S D V E I D V I V K A E Q F G K L V E K T T A A T V T
5598 ttatgttctgaagaattcttcgctaaaagttattgggaattggtgagtacaattatgatattgttacagaagatgaagagtaccct
85 L V P E E S S L K V I G N G E Y N I D I V T E D E E Y P
5682 acattcgaccacttgctcgaagacgtgagtgaaagaaatgctctcactttgaaaagctcgctgttctcaggaatcgccaatc
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5766 aacgattctgcggtatctaaatcaggagcagatggaatttataccggtcttctgtttaaaggcggaagcaattactacagac
141 N D S A V S K S G A D G I Y T G P L L K G K A I T T D
5850 atcattcgcgatgtatcaaccctatcaaggaagggactagaatgctcattccttacaacctaagtgatatttttagcaagt
169 I I R V C I N P I K E K G L E M L I P Y N L M S I L A S
5934 attcctgatgagaagatgtacttctggcaaatgtacgatactactgtctatattcctcaggttcagtcgaaatttatgaaaa
197 I P D E K M Y F W Q I D D T T V Y I S S A S V E I Y G K
6018 ttgatggaaggtatggaagattatgaagacgtttcacagcttgactcaattgagtttgaagatgatcgcggtatccctacagca
225 L M E G M E D Y E D V S Q L D S I E F E D D A A I P T A
6102 gaaatcctcgagcgtattagacgcgttctactattcacttcagcctttgacaaaggaacgctcgaattcttattcttgaaagac
253 E I L S V L D R L V L F T S A F D K G T V E F L F L K D
6186 cgacttcgaattaaaacttctactagcagttatgaagacatcatgtacgcatctgctggcaagaaagtttcgaagaagaattc
281 R L R I K T S T S S Y E D I M Y A S A G K K V S K K E F
6270 acttgccaccttaacagcttactctgaagaaattgtatcaacgctcaccgaagaaacttactgtcttattggaagcgaa
309 T C H L N S L L L K E I V S T V T E E N F T V S Y G S E
6354 accgcaattaaagatttcatcgaatggtgtcgttctcctagcacttcaagagccggaagaataa 6419
337 T A I K I S S N G V V Y F L A L Q E P E E *
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1 M N L A S K Y R P Q T F E E V V A Q E Y V K E I L N Q
10299 ttacaaaattggcgtatcaaacacggctatctattctgtgtggcgctggaactggtaaaaccactactgctcgaattttcgcg
29 L Q N G A I K H G Y L F C G G A G T G K T T T A R I F A
10383 aaggtatgtaacaaaggacttggctctcctattgaaattgatgctgttcttaaatggggtagaaaattgttcgaacattatt
57 K D V N K G L G S P I E I D A A S N N G V E N V R N I I
10467 gaagattctagatacaagttctatggacagcgagttcaaaagttacatcattgacgaggttcacatgctttcaacggagcattt
85 E D S R Y K S M D S E F K V Y I I D E V H M L S T G A F
10551 aatgcgctgttgaaaacattagaagagccctcatcggaacggtgttcattctatgtactactgacccctcaaaagattcctgac
113 N A L L K T L E E P S S G T V P I L C T T D P Q K I P D
10635 actattctcagtcgagttcaacggtttgacttctacgataattgataatgacgacatcggttaactcaacttcaatttatctcgaa
141 T I L S R V Q R F D F T R I D N D D I V N Q L Q F I I E
10719 agtgaataagaaggagctggttatagttatgagcgtgacgccccttcggtttattgggaaaacttgcaaatggaggaatgcgt
169 S E N E E G A G Y S Y E R D A L S F I G K L A N G M R
10803 gacagatcacaaaggctcgaaaaagtccttattatagtcacatcggtgacatggaagcgtttctaatgcactaggagttccg
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10887 gactacgaacattcgcttcaacttctgaagctattgccaactatgacggtcctcaagtggttagaaaattgttaaatgacttccac
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11055 atttcaatcactcaacttctcgtcattttgaaagtaagctagagcaattctgtgaggttttcaatctcactctatttggtg
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11139 atgctagaagaaatgaatgaacttgctggagttgttaaatgggagcctaattgctaaaccgataattgaaacaaacttcttttg
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11223 atgagcaaggaggagtga 11240
337 M S K E E *
dp1ORF014
50961 atgaaagtataatggtcttcaaatgaagcagactcctgaacaaataattgaaaaactttcgagacaacttgaagacgaaggaaca
1 M K V N G L Q I E A T P E Q I I E K L S R Q L E D E G T
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57 S C G M S R N P S Y S G S K V T E A G T V H C F T C G Y
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51297 ttggaacatctagcgaagttagtgcaagcgctcagccctgaagcgtttcgaagaaatgggagaactgaaagtcgagcat
113 F G T S S E V V R Q G V S P E A F R R N G R T K V E H
51381 aaaaatcctcgaagaggaacttgataaatccggtttattcatccttatatgatgaacggaatttgacggaacgagctcatc
141 K I I P E E E L D K Y R F I H P Y M Y E R K L - T D - 4 I
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51549 aacgctcgaagtgttcttcaagttcaccagtcaggtgaagatgacctaaaacggaatttctttatggccaatgatgagctt
197 N R R S V R S K F H Q Y G E D D P K T E F L Y G Q Y E L
51633 gttagcatttccagactattttgaaaaactattagcaagttatcgtagctgagttatcaactgttgactcttgggtca
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51717 atgaagattccagcagtcgctcttatgggagtaggtggaggaaatcaaatcaatttactaaaacgacttcttatagaatatt

253 M K I P A V A L M G V G G G N Q I N L L K R L P Y R N I
51801 gtcttagcacttgaccctgataacgctgggcagacagcgcaggaaaaactctaccgcagctaaagcgaagcaaggtcgtaga
281 V L A L D P D N A G Q T A Q E K L Y R Q L K R S K V V R

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309 F L N Y P K E F Y D N K W D I N D H P E L L N F N D L V
51969 ttgtag 51974
337 L *

dp1ORF015

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113 P Q I S W D N Y L Y M R E R M V E K D K L L P I F H M G
4213gaagactttaaatgggtcaactgattgctcgaaactacattcgaaaggcggaagcatattctctacatttgaattttcaccagcc
141 E D F K W L N L M L G L E T T F E G G K H I P Y I G I S P A
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169 N D S T T K H K K D K W M E R V P E V I R N S S N P D V K
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4549ccaaaacgggttcaagttgaaattccattatcgacgaagaactggagcgcattttagcctagagcaatttagttvgagactat
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309 R L F *

dp1ORF016

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225 E S W Y Y F N R D G S M V T G W I K Y Y D N W Y Y C D A
44169accaacggcgacatgaaatcgaatcggtttatccgtttataacgacggctggtatctactattacgggacggacgtctggcagat
253 T N G D M K S N A F I R Y N D G G W Y L L P D G R L A D
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281 K P Q P T V E P D G L I T A K V *

dp1ORF017

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372

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253 C L R K V S K K G S N A R V C V N E F I R R V K Q V E *
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36183 ggaaaactttagacgttcaagcctttaaagatactccctttagttaaattagggttcagttcaaatgcttacgagttac
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36267 agcgactcaactgttcgaaggtttataagtttcaaccgctttggaggcgatagcttaccaaccaggaaacctactcga
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36351 caatttagagtagaaataagaactacttctcaaatcaaggatattttcgaattggcgaaaaagttcaggacagtttggtag
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36519 gcaaatcaagcgactaacttatttagatacattaaacaggcgacattcttcaagattcctaattggaaaattcaacaattaccatt
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29 F G P T I Q G E G M V I G Q K T I F I R T G G C D Y H C
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57 N W C D S A F T W N G T T E P E Y I T G K E A A S R I L
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2368 gatagaatgaatgatgaaaaccttgactggtcatttaaaatcggttatctttgacgaaaatgacctagcttattgcgcgtgatag
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253 L H T L V Y D N K R G V *
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29 Q S T M F D L Y R N F I H L F M I I K E E Y K M K I E
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85 T E A A V Q R L F G L L G E D A E R D G L Q D T P F R F

373

2840 gttaaagcactcgtgaacataccgttaggtatcgagaagaccctaaacttcctcgaaaaaacattcgacgtcgaccatgaa
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253 A R A E L L Q L I K K *

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113 A S N M K P F R M N I Y V P N Y V G D S I V N Y V K I T
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29 L T R V I E R T Q P E Y N P S T Y Y K P S G V G G C I R
5687 aaaatgtatttcgaaagaatcggtagtctattatagataaacgcagatttcaactaattgcaatggcggaagctggaacattt
57 K M Y F E R I G E S I I D N A D S N L I A M G E A G T F
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6839 caactttcattcttctgtgacgagtagttcgatatcaaggcgaagctctacatttttagagattaagactgaacccatgttcaag
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197 F L Y E N R D N F E K K A Y T F H I T D E M K N Q V L G
7091 aaaattatgacctgcgaagatgtgtagagaaagcgaaagtcctaaaatctattgctcttcagcctattgcccattgttaga
225 K I M T C E E Y V E K G E S P K I Y C S S A Y C P Y C R
7175 aaaggaaggtcgaaatctgtga 7195
253 K E G R N L *

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1 M N A V D G Q V V H I L Q V L A E D G N A T A E K F E K
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29 E V R A A S L V F S R R A A E A V V K G E I Y K D G K N
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57 L S K R V W S S A A R A G N D V Q Q I V T Q G L A S G M
26244 tctgctacagataggctaaaatgctcgagaaatattcgaccctaagggttcgaaaagattgggactttgataagatagctgag
85 S A T D M A K M L E K Y I D P K V R K D W D F D K I A E
26328 aagctagggaaacctgctgctcataaataatcaaaatctcgaatacaatgccccttcgacttgctcgaactaccattagccattcc
113 K L G K P A A H K Y Q N L E Y N A L R L A R T T I S H S
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141 A T A G V R Q W G K V N P Y A R K V Q W H S V H A P G R
26496 acgtgtcaagcgtgtatcgatttagatggtagaatttcttatcgagaagatgctcttcgaccatcctggaatggagtgctac
169 T C Q A C I D L D G E V F P I E E C P F D H P N G M C Y
26580 caaactgtatggtacgaaaactcactcgaagaaatcgctgagtgatgagaggtgggtgagacggagaaactcaatgattgatta
197 Q T V W Y E N S L E E I A D E L R G W V D G E E P N D V L
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225 D E W Y D D L S S G K V E K Y S D L D F V K S Y *

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29 K A I A Y R K V T V K W L P N T D E I Q V Y F D L Y I N
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57 K N R L T M L G T I D P D K S Y F E G I R I V C K K P Q
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18274 tttagagttcattatgtctagccagcactagagcactgttttgcgttgcgttaatataggtgagttttccaagaattggcgg
169 L E F I M S S Q H T R A L V L R C A N I G E F S K N W R
18190 aaatggcaaaaagctatccagctcctgctcgactgtccaaggcggatgactttaagtagacgaaactgtttgggacttttca
197 K W Q K A I Q L L L D Y A K A D D F K V D E T V W D P S
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18026
225 P G S K A G K V A R R K G Y E A I Q Q A L E Q I N K *
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29 R V N K D Q F V E Y D Y K G I K M T I K E R D A K N M K L
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169 Q E R A N G M L P E E V R Y R L Q I E R E K I T L L R A
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197 K M G D Q E I E G E V K D N F V E A L D K A A Q A V W Q
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225 E F S D A T G S Y I K G V T D N D N K P E K *
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57 R R R Y I N C N A I G E D G E T V H P D N C P L C Q N G
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85 F P R I E K L F L Q L Y N H D T G K V E T W D R G R Q S Y
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113 V Q K I V T F I N K Y G S L V T Q P F E I I R S G A K G
53182 gaccaacgaactacttatgaattccttccagagcgttcgggaagacagtgctactcttgaaattttccagaaaagagcgaact
141 D Q R T T Y E F L P E R P E D S A T L E D F P E K S E L
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169 L G T L I L D L D E D Q M F D V V D G K F T L Q E E R S
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225 E R T P S V S R R T P P T R G R G F *
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29 I L A D E K K A D L E S L E D G G E L H L S A S T L E R
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44847 cctgctcgaaaaggttagagtcgttcccaaacctaaaaaagaagtccttgaggagaagaaattcctgaagtttaaggaacagccgga
85 P A R K G R V V P K P K K E V L E E E I P E V K E Q P E
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113 E V G S V S E K S T V R K P A P K K E S V M A I T K A L
45015 gaaagtcgaattgttgaagccttctcgtgctactcgaatcgctcactcagtcctacatcgctcctcaagaagaacttc
141 E S R I V E A F P A S T R I V T Q S Y I A Y R S K K N F
45099 gttactatcgaagaactcgaagaaggtgtttctatttgagttcgcgcaaaaagggttgacagaagacaaaagaacacacttgca
169 V T I E E T R K G V S I G V R A K G L T E D Q K K L L A
45183 tctattgctcctcatctacgaatggcgagttgacggaatttttaaaactcgtcaaggaagaagatattgacaccgcaatggaa
197 S I A P A S Y E W A I D G I F K L V K E E D I D T A M E
45267 ttgattgaagcttctcacctttcttcgctatga 45299
225 L I E A S H L S S L *
dp1ORF029

375

662 atgaaatcagtagttttattatccggcgagtcgactcagccacttggttagcaattgaagttgacaagtggggttctaaaaat
1 M K S V V L L S G G V D S A T C L A I E V D K W G S K N
746 gttcatgctatagcattcaattacggacaaaagcatgaagcagaacttgaaaatgctgctaattgtgcaatgttctacggagtc
29 V H A I A F N Y G Q K H E A E L E N A A N V A M F Y G V
830 aagttccaccattcttgaaattgactcgaaaatctactcaagctctagctcttcttattacaaggaaaagcgaaaatttcacat
57 K F T I L E I D S K I Y S S S S S S L L Q G K G E I S H
914 ggaaaaatcttacgctgaaatcctagcagagaaggaagtagttgacacctatgttccatttagaaatggactaatgttccacag
85 G K S Y A E I L A E K E V V D T Y V P F R N G L M L S Q
998 gctgcggcttatgcttattcgggtggagcttcttactcgtatattggtgctcagcgagacgatcggtggaggtgcttaccct
113 A A A Y A Y S V G A S Y V V Y G A H A D D A A G G A Y P
1082 gattgactcctgagttctataattcaatgtcaaatggaatggaactggaggcaaggttaacctgtcgtcctctta
141 D C T P E F Y N S M S N A M E Y G T G G K V T L V A P L
1166 cttactctaaccaggcgcaagtcgttaaatgggaattgatttagatgttcttatttcttgaactcgttcatgttatgaaagt
169 L T L T K A Q V V K W G I D L D V P Y F L T R S C Y E S
1250 gacgctgaaagtgtggaacttcgcaacttgatcgaccgcaaaaagcgattcgaaagaaatggaatgactgacctattcat
197 D A E S C G T C A T C I D R K K A F E E N G M T D P I H
1334 tataaggagaattga 1348
225 Y K E N *

dp10RF030

20088 atgaataacgaaaaaattattgaaaaaattaaaaatcttattcaattagcaaatgacaacccgagtgacgaagaggggcaact
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20004 gcccttcttatggctcaaaagtgtatgctaaagaataatcgcaacttgctcaagttgaacaatttgatgaacctaaacagttc
29 A L L M A Q K L M L K N N I A L A Q V E Q F D E P K Q F
19920 gagacttctcaagctgttgggaagaagcaggtcgaaatttttggtgggaacgtgaacttggtcatattctcgcgactaatttt
57 E T S Q A V G K E A G R I F W W E R E L G H I L A T N F
19836 aggtgcttttattaatcagcgtgatatcgcttgaataaaagtgaataattttctcggcgaaaaacaagacgctgaatta
85 R C F C I N Q R D M R L N K S R I I F F G E K Q D A E L
19752 gtgctcaaaatatatgaggctgttcttcttcttaccgtattgaccgacttctactcgcgaacccctctacaagaat
113 V S K I Y E A A L L Y L R Y R I D R L P T R E P S Y K N
19668 tcataacctcaaaaggtttttgtcagccttagcctcgatttaaaaagcaggtggaagaatattcacttatggctcctacctagc
141 S Y L K G F L S A L A I R F K K Q V E E Y S L M V L P S
19584 gagcaaaaaaaatgcgcttcaggacacatttcgaaatttaagaaggaaggaattgacagacctcaacatgacttcaatctt
169 E Q T K N A L Q D T F R N L K K E G I D R P Q H D F N L
19500 gaagcgatatattgaaggcggtttcatggcgagaatgcaagattatgccgatgaaatttttggaaaggcggttaactaa 19423
197 E A Y I E G R F H G E N A K I M P D R I L E G G N *

dp10RF031

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1 M A Y Q L E D L L K G L D E P T I K Q V K E I I S K T S
27027 aaagaactcgatgctaaaatttttattgacggcgagcggtcaacattttgtacctcacgcagcttctgtagaagttgttcaacag
29 K E L D A K I F I D G D G Q H F V P H A R F D E V V T C Q G
27111 cgcgatgcagctaacggctcaattaattcttataaagaacaagtcgagcgcttcttaaacaggtcaagataaacggtgatgcg
57 R D A A N G S I N S Y K E Q V A T L S K Q V K D N G D A
27195 cagaccactatccaaaacctcaagagcaactcgacaagcagctctcaacttgcaaaagcgctgtgattacttcaacttctcat
85 Q T T I Q N L Q E Q L D K Q S Q L A K G A V I T S A L H
27279 ccggttgatttagtactcctattgtcccagcagcagacattcttggtatttagaaccttgacaacattacggtcgaaagtgcggt
113 P L I S D S I A P A A D I L G F M N L D N I T V E S D G
27363 aaagttaaaggtcttgatgaagagttgaaagctgttcgtgagctctgtaataactattcaagaagtcgaagttcccgagaa
141 K V K G L D E E L K A V R E S R K Y L F K E V E V P A E
27447 caagaggtcgaagctcagcgagcgagctggaatttaggaatccaggtcgtgctcggtggtggttcccgaaacctcgt
169 Q E A Q A K S P A G T G N L G N P G R V G G G V P E P R
27531 gaaatcggtctcttttgtaagcaacttgctgctgctcaacaaacggcagagcacaagaacaatcatcattctttaataa
27611
197 E I G S F G K Q L A A A Q Q T A G A Q E Q S S F F K *

dp10RF032

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52117 gaccttgatgtcaattgcgtctgcttatcatcggtattttgggatgcttattcctatgcaaaaaggtttaaagcttatct
29 D P D M S I A S A Y H R Y F G M L Y S Y A K R F K C L S
52201 cgacatgacattgaaagcattgcattcgagactattcaaaatggttggcaacggtcaaatcaaaccaaggggccaagttttca
57 R H D I E S I A F E T I S K C L A T F K S N Q G A K F S
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85 T Y L T R L F K N R I V L E Y R Y L N A P S M N R N W Y
52369 gtagaagtgacgttcgatagcgtttcgacaaaatgaagaaggcgacgatttagtatcctatcgacagttggctattgtgaagac
113 V E V T F D S V S T N E E G D D F S I L S T V G Y C E D
52453 tacggaaaaattgaaattgaagcaagcttgcattcagcgttttctaatacagagtagtcttatctcgtctgctcattcaa
141 Y G K I E I E A S L D F M T L S N T E Y A I S V I Q
52537 aacgctcttcagtaagcgacgcagaaattgcgctgaaattggagtaagcaggtctgctattagtcagtcgaagtcacta
169 N G P S V S D A E I A R E I G V S R S A I S Q S K - K - S L
52621 aaaaataaataaagattttatataa 52647
197 K N K L K D F I *

dp10RF033

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376

29 I I N K V V D E I V E A A C G S L D Q A M E E I Q I V V
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57 S Q N P V I M E D L N Y Y I G Y L P T L L Y F A A D R A
7922 gaaatgggtggaatacaaatggattcaagttctgtcatcaggaaagaaaatacagataatctatacattttgacggcggggaaa
85 E M V G I Q M D S S S A I R K E K Y D N L Y I L A A G K
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113 T I P D K Q A E T R K L V M N E E V I E N A Y K R A Y K
8090 aaagttcaattaaagctagaacaggccgataaggtatttagcatctttaaacaagaattcaaacctggcaactagcagatttagaa
141 K V Q L K L E Q A D K V L A S L K R I Q T W Q L A E L E
8174 actcagtcataataattcaaaaggagttattataatgcaaaagacgtagacgtgaaaatgattga 8239
169 T Q S N N S K G V L L N A K R R R R E N D *

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1 M S Q N T T R T D A E L T G V T L L G N Q D T K Y D Y D
215 tataatccagacgtctcttgaactcttccctaacaacatccctgaaaataattacctagtagaacatttgaggatattgaattcact
29 Y N P D V L E T F P N K H P E N N Y L V T F D G Y F T
299 tccctttgcccataaacaggacagcctgacttcggaattgttttcattagttacattccaaacgaaaagatgggtgaattctaaa
57 S L C P K T G Q P D F A N V F I S Y I P N E K M V E S K
383 tcattgaaattgtactattcagtttccgtaaccacgtgacttccacgaagattgcatgaacattatttgaatgactttgat
85 S L K L Y L F S F R N H G D F H E D C M N I I L N D L Y
467 gaatttggaacctaagtagcattgaagtcagggccttactcctcgtggtggaattttcaatttaccattcgtcaacaaa
113 E L M E P K Y I E V M G L F T P R G G I S I Y P F V N K
551 gtgaattcctcaatttgcaactcctgaactgaacagcttcaacttcaacgcaaatgaacttcttggaatgttcaaggtctt
141 V N P Q F A T P E L E Q L Q L Q R K L N F L G N V Q G L
635 ggacgagctattcgatag 652
169 G R A I R *

dp1ORF035
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1 M H L M K D S K M L R T W K S L A F E F E T K V R T T S
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29 G L K L S P A M K T M T R T K I W K G Y K M K V F I N N
17257 catactgaagctgatattgactacaagatattctaaattttgtagcttatcgaaactctcctaaccctcaaatccaaatcact
57 H T E A D I D Y K D I L N F V A Y R N S P N P Q I Q I T
17173 agctggaacgctttgctttcctgctatacaggaatgagctttcttataaaggagtttcaataacggacttttttgaagcatt
85 S W N A L L S C Y T R N E L S Y K G V S I T D F F E A I
17089 caaactattgcaagttccttactcacctagactcgaaaacaaattgatacacaataatgaaaagcgactcgaaaggattgaggaa
113 Q T I A S S F T H L D S K T I D T Q N E K R L E R I E E
17005 cttcagtcagaataggtcattgtaactgtactatcgacgaacttaaaaaaggagttccacgaaatgccggatattgaatcagct
141 L Q S R I G H C N C T I D E L K K G V H E M P D I E S A
16921 atttcttaccagtagcgacagattttgttattgaagatgaacttaattttctgctaaactaa 16859
169 I S Y Q Y G Q I L A Y E D E L N F L L N *

dp1ORF036
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1 V L V E R K A D K E C W E W L E A V R A N I V E E V R N
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29 G L S I V I A S N T V G N G K T S W A V R L L Q R Y L A
48976 gaaactgcacttgacggaagaattgttgagaaggaatgtttagtagtgcagctcaactattgactgagttcggcgactataat
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49060 tattttcaaacatgcaagaatttctcgaacgtttcgagcgcttaagacttgtagctattagtcagacgaataggtgga
85 Y F Q T M Q E F L E R F E R L K T C E L L V I D E I G G
49144 gggtcccttaaccaaggcctcttactccttactgtgacttggttaattatagggttgacaataacatttgcactatttatagc
113 G S L T K A S Y P Y L Y D L V N Y R V D N N L S T I Y T
49228 actaattatactgacgatgaaattattgaccttttagggccaaaggctttatagtcgtatataatgatactcagtggttctagat
141 T N Y T D D E I I D L L G Q R L Y S R I Y D T S V V L D
49312 tttcaggcaagcaatgtaagaggttgaggtaagcgaaattgaatcatag 49362
169 F Q A S N V R G L E V S E I E S *

dp1ORF037
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1 M V K L K S K I Y S V A Y I I L V V I A N L V T I Y F
55939 gaaccttttaattgtaaggaattttaattctccaagcagttgggtttatgggattcactttcctgcttataaatctaataagc
29 E P L N V K G I L I P P S S W F M G F T F L L I N L I S
56023 aagtcagagaagccaaaatttgaggttctttgatagtggttattccttacctcgttgatttcttatgcaaaacctta
57 K Y E K P K F A G S L I W V G L F L T S L I C F M Q N L
56107 ccacaatcgcttgctggtcctcaggagttgcatcttttgataagtcataaagcaagtgcttttatattcgacaagctctcgaat
85 P Q S L V V A S G V A F W I S Q K A S V F I P D K L S N
56191 aaatttagactcgaagattgcaaatgctttgtctagcaacatcggttctattatagacgcaaccatattgatttcattaggactg
113 K L D S K I A N A L S N I G S I I D A T I W I S L G L
56275 agtccctttggaattggaacggttcataatagatattccgctcagcgtactagggccaaagttctagttcagtttatcttgag
141 S P L G I G T V A Y I D I P S A V L G Q V L V Q F I L Q
56359 tcaattgcttcgagatatttgaaaaagtag 56388
169 S I A S R Y L K K *

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1434 acttacaagtcgaaatttcttagcagcggaacttatgaccacggttcgagtcgaagggatgggttggtgacttttatcacgtc
29 T Y K V E I S L A G G T Y D H G S S Q G M V V D P Y H V

377

1518 aagaaaatcgacggtacattcattgacagacttgaccacgctgttcttctcaagggaatgaaccaatcgcttttagcaaatgca
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85 V D T K R V L F G F R T T A E N M S R F L T W T L T E L
1686 atgtggaagcatgctcgatcgactctatcaaaactatgggaaactcctacaggttgccgagaatgtacttactacagagattttc
113 M W K H A R I D S I K L W E T P T G C A E C T Y Y E I F
1770 acagaagacgagattgaaatgttcaagaacgtaacctttatcgacaaagacgaaaagattactgtccgcgaaatttttagagcag
141 T E D E I E M F K N V T F I D K D E K I T V R E I L E Q
1854 gagcaggataatggttaa 1871
169 E Q D N G *
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1 M N K S A T F W L V R T A L I A A L Y V T L T V A F S A
3390 attagttatggacctattcaatttagagtcagtgaaagccttgattcttctacctttatggaaccatagatggactccggggatt
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3474 gtattaggaacaattattgcaaaacttctttcactcttgactgattgacgttttattcggttactgtacttcttcttggga
57 V L G T I I A N F F S P L G L I D V L F G S L A T F L G
3558 gtagtggcaatggtgaaagtgtcaagatggcaagtcctctatattcacttattctgtccagttcttctgtaattgcttaccttatt
85 V V A M V K V A K M A S P L Y S L I C P V L A N A Y L I

3642 gcgctggaacttgaatagtttactctttaccttttgggaatctgtcatctatgttaggaattagtgaagcgattatcgtttta
113 A L E L R I V Y S L P F W E S V I Y V G I S E A I I V L
3726 atttcatacttcttatttccacgctggcgaagaacaatcattttagaacactgataggagcgaataatgggatttaa 3803
141 I S Y F L I S T L A K N N H F R T L I G A K N G I *
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29 Y D T T N G F R G V A N P C D Y I A A T N F G T L F I E
7360 ctgaaaactactaaagaagcttcttgcagcttataacatcactgataatcaatgggtccagctatcacgcgcagatggatgc
57 L K T T K E A S L S F N N I T D N Q W F Q L S R A D G C
7444 aaatttattctcgcggaatttttagtgatttccaaaagcatgaaaagattatggtatccaatttcaagccttgaaaaaatt
85 K F I L A G I L V Y F Q K H E K I I W Y P I S S L E K I
7528 aaacggctctggagttaaaagcgtcaacccaaacttcatcgatgaggggtatgaagtttcttacaagaagcgtcgaactgattg
113 K R S G V K S V N P N F I D A G Y E V S Y K K R R T R L
7612 accattccttccaaaatgttctagatgcagttgagttcattacaaggagaaaagcaatggcaagacttaa 7683
141 T I P F Q N V L D A V E L H Y K E K S N G K T *
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8208 atgcaaaaagcgtagacgtgaaaatgattgaccctaaacttgaccgattaaaatacacaggtgattgggttgatgtacgaatt
1 M Q K D V D V K M I D P K L D R L K Y T G D W V D V R I
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29 S I T K I D A D S A D V S R C R K V L Q K A V Y S V
8376 gcggcaggtgaatgcattaaaattgcacacggatttgccttgaacttctcaagggtatgaagcaattctgcatcctcgttcc
57 A A G E C I K I A H G F A L E L P K G Y E A I L H P R S
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85 S L P K K T G L I F V S S G V I D E G Y K G D T D E W F
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113 S V W Y A T R D A D I F Y D Q R I A Q F R I Q K P A
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141 I K F N F V E S L G N A A R G G H G S T G D F *
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48334 aactgggttcaacttttcgagcactatgaaaaatgttcgaacttatttaaacattgagtcgaacattgcaacttttgcgatttta
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113 A E S F V K Y E N V R K R L N L S E R F I T V S T F K R
48502 gcctggatttggacgaactcgaaggaaaaacgggttcaaaattcgaaggattttattag 48561
141 A W I L D E L E G K T G S K F E G F Y *
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29 P I H V K I R A A G V M N L I A N G K I P N T L G K V
31867 acagaactgtttggagaaacttcgacagtcactaaagacaatgctagtctagcatcaattactgaccaacagaagaaagagcg
57 T E L F G E T S T V T K D N A S L A S I T D Q Q K _ K _ E A
31951 ctgaccgattgaacaaaacgataccggtattcgaagacatggctgaacttcttcgagttattcgcagaagettcaatggtagag
85 L D R L N K T D T G I Q D M A E L L R V F A E A S M V E
32035 cctacttacgctgaagtgcggagtatatgacagatgagcaacttatgacaattcttcagtgcaatgtacgggtgaagtgaactcaa
113 P T Y A E V G E Y M T D E Q L M T I F S A M Y G E V T Q
32119 gctgaaacctttcgtacagacgaaggaaatgtctaa 32154
141 A E T F R T D E G N V *

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25498 ttcgatatgtttcgaaataatctgttttagatgtaaggtcgaaacttatgctcacaatgggcacaattaaccttgaaactgtgggt
57 F D M F R N N L F R C K V E L M L T M V T I N L E R L G
25414 cgactccttctcggttggtgttcagttgtttttttttgtcatcaacttcgtcttcttctcactcgtttctctgtgaggt
85 R L L L R L V V Q F V L F L C H Q L R L L H S F H L B A
25330 cctcttgttctgttaattcgtttgttaatacaggaatgctccagctgagatttcgtcaagctgagcaagttcttccaaatgc
113 P L V R L I R L I Q A M L Q L R F R Q A E Q V L P K C
25246 gttcccatctctgttcgccccttttcttctactga 25211
141 V P I P C P P P P S Y *

dp1ORF045
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29 F K V N C D H C E H K F D L T S K Q I I S K H I E K G V
25508 gagggtgattcttcgaatgtcctaagtgccattacgggtccaccatttatgtaggaaacaggaaattgaaacatttattcga
57 E W R F F E C P K C H Y R F T T Y V G N K E I E N L I R
25592 tttagaaatacttgcagctaaaaatgaagcaggaaacttcaaaaggagctgctgtaatacaaacacttaccattcatatcga
85 L F E W K S N K A K S V L E D I S T T L S T L K Q Q V D
25676 attcaggatgagcaagctgggcataaaatctcagggttatggcgaagctaaagaaggagataaacattgaaacgagaaaaa
113 I Q D E Q A G H K I S G L M A K L K K E I N I E K R E K
25760 gaatgggtatctatatag 25777
141 E W V S I *

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57 G I D Q T T V A I N H Q N D V I Q D G T R K I Q R Y R L
43026 tatcagcacttaaaagggaagtgataacaggctatacaactctcgaccatttttagagagctctctattttattcgaaagttat
85 Y H D L K R E V I T G Y T T L D H F R E L S I L F E S Y
43110 aagaaccttggcggaatggtgaagtgaaagccttgatgaaaaatacaagaattaccgaattgggaggaagatttagatgaa
113 K N L G G N G E V E A L Y E K Y K K L P I R E E D L D E
43194 actatctaa 43202
141 T I *

dp1ORF047
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29 Q V K S L R D A L K E Y M K E N D I E S A Q G K H F S A
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85 E M C E K L S G L I E Y K P V I N T K L L E D M I T
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47961
113 G E I D Q E A I L P A V V I S V T E G I R F G K A K I *

dp1ORF048
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16457 agaattgaaatcatcaagaatataactacaagaatcgaacgccttaacgaagaattaaagcaagaatgaacaaggtaacaa
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16373 gaaagccgacactagtagtctgcgctagaagattgcgctcgtcaaattgctggaatttatcaataa 16308
113 E S R H L V S A L E D C A R Q I A G I Y Q *

dp1ORF049
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43766 tccgtcgtaggcgtgattgcagtgatgatgttatcactgtcaatgaacatccctgtgatgacctccagcgcctcggttagcacc
85 S V V G V V A V N D V I T V N E H P C M T S S A C A S T
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113 F A S P D E D V A S F S I P R S I F T N *

dp1ORF050
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379

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15249 cctgaaagccaacgaatgtcttgaggctatgtgtattagatgacctccagtcactaatgcggccgctgaaattggataccac
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29 S W V S D G Y G G K K D K A N E V V A D D L V C L V D
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30017 tatgatgaaggcaaaatcattcaacgagccgatactatcgaaattaaaaactcaggaagacggtacagggtagtagaaaccac
85 Y D E G K I I Q R A D T I E I K N S G R R Y R V V E T H
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113 N L L E Q D I L I E L K L E V N D *
dp1ORF052
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29 V E F D E Q D T D R P D D Y I V L R Y S H R M P S A T N
30684 agcctgaaggttttcttattggaaagttaaactctacgtccattcaactcaattattggatcgacgaatatacgagaag
57 S L G S F A Y W K V Q I Y V H S N S I I G I D E Y S R K
30768 gttcgaacattatcaaggacatgggctacgaagtaacctatgcagaactggtgactctcgacacaaatgctttctagatac
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30852 cgactagaatcgaaatatagaattccacaaggaggaaactaa 30893
113 R L E I E Y R I P Q G G N *
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113 S S F G I I F A I A M L L S T *
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29 A A Q I P A T A A T Q V G N K K Y I L A G T C V K N A T
27795 acatttgaaggacgcaaaactggactcgaagtagtattaccggtgaacaatttcgacggagttatcttcgctgaccaagaagt
57 T F E G R K T G L E V V S T G E Q F D G V I F A D Q E V
27879 tttgaaggtgaagaaaagtaacctgacagatttagttcacggattcgtcaaatatgcagcccttcgaaaagttggcgatgct
85 F E G E E K V T V T V L V H G F V K Y A A L R K V G D A
27963 gtgctgaatctaaaaacgcaatgattcttgcgttaaatag 28004
113 V P E S K N A M I L V V K *
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29 V P G T P Y R L Q V W V K M S L V K I E T R A G N G Y Y
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57 K R L V C Q D D F V F Y G K E S I D G Y L I D A T T I T G
18899 aaatcttggcggaattattgtgagcctatgaacagggcatattctcgaaactattgcatcgagagaagcagctgaactgaacaga
85 K S L A E Y C E P M N R H I L E T I A S R E A A E L N R
18815 gctaaaaagcaagacacagaatggagatactag 18780
113 A K K Q D Q Q K W R Y *
dp1ORF057

381

29108 atggctacattgaaagctcttagcaccttaacgtttccggagcagtagtcattcagggtcggtattttcttgccttgaagcg
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29 L A S S L I E R N F A F E I K A A E D G E T V E T V P Q
29276 acaattgaatcagttgaagaaattgacgaagtgaacaaatgcgcgaagatgcggctaaaaccgttccctgagctcggtgaa
57 T I E S V E E I D E V E Q M R E E Y A A K T V P E L V E
29360 ttagcaagagctaattggaattgacatttctcaatttctcgaaaaagcgaatatatcgacgctttaattaagtacgaactagga
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29444 gaggtaa 29449
113 E *

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29 V F I K F L R N D F M L D F F S S P I S S K R F R A D
51329 gccttcgctaactcttcgctagatgttccaaaattctctttcagccactgggttccatagaacctccatcgcttcgacctaa
51246
57 A L P N Y F A R C S K I P F Q P L V S I E P S I V S T *

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1 V T N C V R W K Q Y H F T V V N Q V E L T N V T N V R K
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29 F V S V S E L S N F L R V D S D L K T C F F S D E F L S
28730 gtcacttgcaagaagcaagaagttttcccaagaaccttgaaacacaaatgcaagagctttcttgatagagtcactcttagtcat
57 V T C K K Q E V F P R T L N T N C K S F L D R V T L S H
28646 ttgggtataaagtgttccggttcaagaccattcgagtagggcgaaacacctgtacgattttcgatgtcatccattgctgctaa
28566
85 L V I S V S V Q D H S S R A N T C T I F D V I H C C *

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29 G A G F R T V D F S L T E P T S S G C S L T S G I S S S
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44809 gcagcttcttcttttttaggtttagtttcatcttccattgtgtaccaacggttcgagagttgaagctgaaaggtga 44735
85 A A S S F L G S V S S S I V Y Q R S R V E A E R *

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1 M A A Q T D I E L V K I N I D N D N S P S P M T D Q S I
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29 S A L L D K H K S V A Y V S Y M I C L M K T R N D V V T
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57 L G P I S L K G D A D Y W K Q M A Q F Y Y D Q Y K Q E Q
29703 cttgaaactgatgaaaagtcgaacgctggttcgacaatttcaatgaaaagggtgatgggacatga 29768
85 L E T D E K S N A G T S I L M K R A D G T *

dp1ORF069
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29 L A E S Y E K A L A F L S L R N V D T I V V L E L E V D

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57 I E K C T E S F D H N E K M F C S L F H F D T C R A W T
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85 Y D K T I E V D D I D F S K A R K Y D R K *

dp1ORF070
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29 P I Q E D D I Q F V D I T G L D P I V R E N V L E L I S
16141 cggagccgtgtaggagtttcaaatatggtacaaacctcgaccagaatgatgtcgacgatttctacagcagccaaagaagaa
57 R S R V G V S K Y G T N L D Q N D V D D F L Q H A K E E
16225 gcgcgtcagctttgctaactacctaaccaagctacaaagtcaacaaaagcaaaataaatag 16284
85 A L D F A N Y L T K L Q S Q Q K Q N K *

dp1ORF071
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29 V R N I L T S L S L I V Q T V R D L V I L T A D E H T S
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57 V S I K I S I P S I Q K T L Q P I H G R N G R G M T E L
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85 K G Y P G S Q A Q T V R L I I S I *

dp1ORF072
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29 S K C F N S F P C N L T S K T S S R P R G F C F Q W R A
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57 F A F F S S F F A F L F E S Y K S I G S S F N V P H I F
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85 D D F S V F A I S V F N D R *
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29 K Q L Q N L L E K L Q R L L V A L A L K R K V E I K C V
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57 K I V K T K H S I L E F S M K M K V A M S T P H S L T R
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85 R F A T P Q Q L L A I E R *
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29 L H G F W V N C S K N D F G Y L K L H K S I K S C S K S
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57 S A T A R T R V F E V L S N W F C F N R I R E R T Y D C
32550 ggttacccttctcttattgggtattgcagcgctctatttaa 32591
85 G Y P S S Y G I C S R L Y *
dp1ORF075
22447 atggcaagttttgtcctgtaattccgtcatggcccaagggaatgaaagagccatcgatactgttttctgaaacgaatg
1 M A K F C P L N S V M A Q R E N E R A I D T V F P E R M
22363 gaaccgctcgtatgacgatatcgaaagttcgaaaaggtgagccctttgtccaccatgttaggagctggagttgttcttacta
29 E P S A M T I S K V R K G E P F V H H V R S W S C F L L
22279 aaagggcgaagttgaacttaggtattttctcaggttattgtcattatcagtcactcctttaaaggaacgtgttgc
57 K G T K L N L G S L F L R L I V I I S H S F N V G T C C
22195 gtcactaaattcttgccaaacggcttgagctgctttatctag 22154
85 V T K F L P N G L S C F I *
dp1ORF076
5728 gtgagagcattttcttctcactcagctcttcgagcaagtggtcgaatgtagggtactcttctcttctgtaacaatatcaatattg
1 V R A F S S L T S S S K W S N V G Y S S S S V T I S I L
5644 tactcaccatttcccaataacttttagcgaagattcttcagggaactaatgtgacggttcggtggttcttctacaagttt
29 Y S P P P I T F S E D S S G T N V T V A A V V F S T S F
5560 ccaaaactgctctgctttcacaatcagctcaatttcaacactcgtgctgataatgcatcgaaggaagtttgagccatcagct
57 P N C S A F T I T S I S T S L S I M H R R K F E P S Y A
5476 gtaaacatgacgattcgcgtcaccaaaaatagccaatag 5435
85 V N M T H S P S P K I C Q *
dp1ORF077
14800 atggaacgaataaagacgctatttcacgtgatttatgctaacggcactcatttagaagtagcagctttgttcgataccggtgat
1 M E R I K T L F H V I Y A N G T H L E V A A L F D T V D
14884 gattatgatgagcttagaggacatccaggggtatattgataccctgacctttataatcaaaggagcatttagaagggcgct
29 D Y D D V I E D I Q G Y I D T P D L Y N Q R S I R M A P
14968 tacaatcctgacatcaatggcgactattgctactgacattttactacgactagatgatattatctacgtcgacgcaactgt
57 Y N P D I N G D A I A T D I L L R L D D I I Y V D A T C
15052 gaaactattaaatcagaggagcctattgcatga 15084
85 E T I K Y E E P I A *
dp1ORF078
17507 atggcaacagtaaaaggaaacagtaaaatttgacggacgtcttgtaactatcttcgactacgacgatttagagtggaaggat
1 M A T V K E T V K F D G R L V T I F D Y D D L E W E G Y
17423 gcacctaataaggattcgaagatgttaggacatggaagtccttagcattcgagttcgaaacgaaggtgaggacgacgagtg
29 A P N E G F E D V E D M E V L S I R V R N E G E D D E W
17339 gttgaagttatcgctgctatgaaaacgatgacgaggacgaagatttgaaggggtataa 17280
57 V E V I A C Y E N D D E D E D L E G L *
dp1ORF079
35288 atggaactgataccattgataaatcctcgaacaaggttgacccctgcgcttaccatttgtccagcaatccagtaaccttagaa
1 M E L I P L I N P R T R L T P A L T I C P A N P V T L E
35204 acaattgaagttcccatgctgccaatttttagagacagctgaaccaatcattgacccaataaccactaataagtttcaatcagg
29 T I E V P M L P I L E T A E P I I D P I P L M K F R I R
35120 ttcgcacctcctgaaacatctgtcccacaaagctagcaatcttgtaactaatgatgaaagcattgttccagctgtcgataaaa
57 F A P P E T I C P T K L A I L L T N D E S M F P A V D K
35036 agtgagccgagaagtgaaacacatttga 35007
85 S E P R S E A I P *
dp1ORF080
42490 atgttgaaaccttacaataatcgcgcaaaattgtggcagagttcactattggacaaggagctgaaagaaacttgcctcaaaacacg
1 M L N L T K S R Q I V A E F T I G Q G A E K K L V K T T
42574 attgtgaacattgatgcaaacgagtatcaacgctctctgaaactcttcagaccagacttgatgctgcgaaccgtcgagaa
29 I V N I D A N A V S T V S E T L H D P D L Y A A N R R E
42658 cttcagctgacgagcaaaaacttcgcgaaaactcgttacgcaatcgaaagatgaaattctagctgaacagtcaaaagactgaaaca
57 L R A D E Q K L R E T R Y A I E D E I L A E Q S K T E T
42742 gctctaacagctgaataa 42759

dp1ORF091
43189 atgaaactatctaacgaacaatatgacgtagcaaaagacgtggtaaccgtagtcgttccagcagcgattgcactaattacaggt
1 M K L S N E Q Y D V A K N V V T V V V P A A I A L I T G
43273 cttggagcgttgatcaattgacactactgctatcacaggaaccattgcacttcttgcaacttttgaggtagctgtcttagga
29 L G A L Y Q F D T T A I T G T I A L L A T F A G T V L G
43357 gtcttagccgaaactaccaaaggaacaagaagctcaaaacaatgagggtggaataa 43413
57 V S S R N Y Q K E Q E A Q N N E V E *

dp1ORF092
46989 atgaaactatctccatattaaggaagacactaaaaggaagccggacaggaacggaagaaaactgcactcgaactagctcaa
1 M K T I S I L R K D T K R K P D R N G R K T A L E L A Q
47073 gagattgatatgtcacctagtgagtttagcagagctccttcaaattcctgaaaggacggcaaccagaattttaaaactcgacaaa
29 E I D M S P S E L A E L L Q I P E R T A T R I L K L D K
47157 ctgctcaacaagagcaatgctcaataatagaagggtatataaatgaaattcactga 47213
57 L L N K E Q C S I I E R Y I N E I H *

dp1ORF093
45756 atgcaacatcagattaaacaatgtttgaaacttgcttctgctcaatgcaatcaattgcctgtttagtttccctaaacct
1 M Q H T I K Q C L K L A F L L T A I S I A C L V F P K P
45672 tgctcatcgctaaaaggaacatggatgctcttgctgctattcgaaacattcaacctgggtgcgcaatggagtagtcttgaaac
29 C S S P K R K H G C S C A Y S K H S T W C A N G V V L N
45588 gaaaactgctcattgcttgaagaagctattcggtttcgagagtcattgtag 45538
57 E N C S L L E E A I R F R E S M *

dp1ORF094
8281 atgtacgaattagttctatcactaaaattgacgccgacagcgccgatgtctcaagatgtcgaaaagtgtctcaaaaggctcaag
1 M Y E L V L S L K L T P T A P M S Q D V E K C F K R L K
8365 tatattcagtgccgaggtgaatgcattaaaattgcacacggatttgctcttgaaacttctaaagggatagaagcaattctgc
29 Y I Q W R Q V N A L K L H T D L L L N F L R D M K Q S C
8449 atcctcgttccagtccttttaagaaaactggtctaa 8484
57 I L V P V F L R K L V *

dp1ORF095
8877 gtgggaaaactacttcagctctcgacattgtcaagaatgcgcaaatggatatttgagcaggaatgggaacagaagactgaagaac
1 V G K L L Q L S T L S R M R K W Y L S R N G N R R L K N
8961 tcaaggaaaagctggaaaatgcgctgcatccaaagctagcaagactgctgtcaaggaaactgaaatgcaactcgatagctcttc
29 S R K S W K M R V H P K L A R L L S R N L K C N S I V F
9045 aagagcctcttaagattgtatatcttgaccttgagaatacattag 9089
57 K S L L R L Y I L T L R I H *

dp1ORF096
46681 gtgattcataaattcttcaatttcggtgaacttatctgcggtttctcctgttaccaggttgcatcttgactgtcttcgaaagtat
1 V I H K F F N F V E L I C G F S C Y Q V A F D C L R K Y
46597 cttagcaagaggttcaataaccttttcccaattgctaaatatcagcaggactttccttgctggatacattcctcgacaatttc
29 L S K R F N N L F P I A K Y H A G L S L L D T F L D N F
46513 gatacatcttgcgaacttgcaagacttgacatcttgatagtagttaa 46469
57 D T S F E L A R L D I L S S *

dp1ORF097
39100 atggacgggattgaaatcttgatactgaccgacgtatgctcgctcgctgtcagtagtactaaatccctcaccgtttggactatt
1 M D G I E I L I L T D V C S S A V S M T K S L T V W T I
39016 agagaaagcaggtgagtagatattgcgaacgtccgtcagctcctgcaggtccaggaattccttgaaagccttgaggacctgaag
29 R E S E V S I L R T S V S C R S R N S L K P L R T L K
38932 accttgaaactccttaggacctgtttcacctatcttggaactga 38888
57 T L N S S R T C F T Y L G N *

dp1ORF098
43627 gtgaaaatgctccgtgggatgctaaacagggcgacatcttcatctggggacgcaaagggtgctagcgcagcgctggaggtcata
1 V K M L R G M L N E A T S S S G D A K V L A Q A L E V I
43711 cagggatgttcattgacagtgataacatcattcactgcaactacgctcagcaggaatttccgtcaacgaccacgagtagcgtt
29 Q G C S L T V I T S F T A T T P T T E F P S T T T M S V
43795 ggtactatgcaggtcaaccttactactacgtctatcgcttga 43836
57 G T M Q V N L T T T S I A *

dp1ORF099
38298 atgcaagttcgccatctgctactgaagctccagctgggtgaggtctacgcaagttcctaccgtcccaggtggtcagtagttat
1 M Q V R H L L L K L Q L V D G L R K F L P S Q V V S I Y
38382 ggactcgaacaagatggcgctacactgacaaactgatgaaattggatattcagtttcaagaatggcgagcaggggtcctaaag
29 G L E Q D G A T L T K L M K L D I Q F Q E W A S R V L K
38466 gtgacgcaggtcgtgacggtattgcaggaagaacggaatag 38507
57 V T Q V V T V L Q E R T E *

dp1ORF100
1597 atgcagttgacaccaagcgagttctatttgatttagaactacggctgagaatatgtcaagattccttacctggactctcacgg
1 M K L T P S E F Y L D L E L R L R I C Q D S L P G L S R
1681 agcttatgtggaagcatgctcgtatcgactctatcaaaacttggaactcctacaggttgccgagagtagtactactacgaga
29 S L C G S M L V S T L S N Y G K L L Q V A Q N V L T T R
1765 ttttcacagaagcagagattgaaatgttcaagaacgtaa 1803
57 F S Q K T R L K C S R T *

dp1ORF101
19220 gtgataatttttagtccagttccactacatttgaagcgcgattaggtcatctaggctgtctagctcgagttcgattacaaggt
1 V I I L V Q F P L H L K A R L G H L G C L A R V R L Q G
19304 tgccagtagtcaatttcacaaaagtaagcgacatttccaactttcttagtgcttcagatacctatcatatgtcgctcttcgt
29 C Q Y Q F H K S K R H F Q L S L V L H D T Y H M S P L R

385

19388 caaatagtcgagcagaataaacttcgaatttcatttttag 19426
57 Q I V A Q N K L R I S F *
dp1ORF102
4034 atgataacgtgggaatgtttgactgtatcgccgaactcgataaaattcctggtgtatttagacagcctaagacacgtgaacagc
1 M I T W E C L T V S P N S I K F L V Y L D S L R H V N S
4118 ttttggagcaccacaaatttctgggataattatctatacatgagcgagcgaatggttgagaaaagacaagctcttacctattt
29 F W K H H K F L G I I I Y T C A S E W L R K T S S Y L F
4202 tccatatgggagaagactttaaatggctcaactga 4237
57 S I W E K T L N G S T *
dp1ORF103
49352 ttgaatcatagatatagtaacatcacactatttttcttggcagattgtctttctttgtattttgctgagcggtgtcctattgt
1 L N H R Y S N I T T I F L W Q I V F L C I C C A V S Y C
49436 gcaggagtgacataatgagcgagagctctcaagataaggtgattcaagttataagcagaagaaaagtcagccgtctacttgaca
29 A G V H N E R E S Q D K V I Q S Y K Q K E K S A V Y L T
49520 gtogatagttcaggagcttggctaggaagtgctccgggagccaaggaaagtcctctctacaatgaaaagggacagcatgtagga
57 V D S S G A W L G S A P G A K E S P L Y N E K G Q H V G
49604 aaattgaaagaggtgggagagtgga 49627
85 K L K E V G E *
dp1ORF104
21427 atgagaaaaagagtgattttgaagctaaaaaggttgaactggtatgtccttaattcctactctcgaatggttgagttttcgaa
1 M R K R V I L K L K R L N W Y V L N S Y S R M V E F F E
21343 cttttgaacttttgcgaatgggttcgacttttgcgaaggttgaggttttcgaaccggttgagtttttcgagcatttcgacttttc
29 L L N F S N G S T F R R I E V F E P V E F F E H S R L F
21259 gacccctttctatgctcgacttttcgagtggtttga 21224
57 D P F L C S T F R V F *
dp1ORF105
2028 atgatagtcgatccaccagttcgaatgaaaatagtccttttgacctataaaccattccttcaccttgaattgtaggaccgaaaaat
1 M I V A S T S S N E N S L L T Y N H S F T L N C R T E N
1944 ttccatgatagcatttttctcagggtcgcaacattgattcgaatcttgcctcttcaggctgattgtattgattaaccattat
29 F H D R H F L R V A N I D S N L A S F R L I V L I N H Y
1860 cctgctcctgctctaaaatttcgagacagtaa 1828
57 P A P A L K F R G Q *
dp1ORF106
10529 atgaacctcgtaatgatgtaactttgaactcgctgtccatagacttgatctagaatcttcaataatgttttgaacattttc
1 M N L V N D V N F E L A V H R L V S R I F N N V S N I F
10445 tacccattattagaagcagcatcaatttcaatagagagccaagtcctttgttcacatccttcgcaaaattcgagcagtagt
29 Y P I I R S S I N F N R A K S F V H I L R E N S S S S
10361 ggttttaccagttccagcgccaccacagaatag 10329
57 G F T S S S A T T E *
dp1ORF107
10750 atgagcgtgacgccctttcgtttattgggaaacttgcaaatggaggaatgcgtgacagtatcacaaggctcgaaaaagtccttg
1 M S V T P F R L L G N L Q M E E C V T V S Q G S K K S L
10834 attatagtcacacgttgacatggaagccgttttcaatgacactag 10878
29 I I V I T L T W K P F L M H *
dp1ORF108
49447 atgcactcctgcacaataggacaccgagcagaatacaagaagacaatctgccaaagaaaaatagttgtgatgttactata
1 M H S C T I G H R A A N T K K D N L P K K N S C D V T I
49363 tctatgattcaatttcgcttacctccactcttacttgcttgcttgaaaaatctagaaccactgaagtatcatatatacagac
29 S M I Q P R L P I L L H C L P E N L E P L K Y H I Y D
49279 tataaagcctttggcctaaaaggtcaataa 49250
57 Y K A F G L K G Q *
dp1ORF109
31632 atgtggtgtgcaagtcaccaatagttgatttctccttcaactttccagcctttgaaagccttacctgttaaggtagggtaact
1 M W L S K S Q I V D S P S T F Q P L K A L P V K V G S T
31548 gggttttgagaaaatcttcttacctgcttcaactcgaactgcgtcgccggttctgttccaccgttcaaatcgaatgtcacgcga
29 G P G E I F L P A S T R T A S A V P V P P F K S N V T R
31464 cgaagaaccgctggaagttgtgccacatag 31435
57 R R T A G S C A T *
dp1ORF110
16444 atgatttcaattctagcatcaacttccatgtcgcgagtaagttgtgactccagtttcagcgacaggacatgctttgaataactgca
1 M I S I L A S T S M S R V S V T P V S A T G H A L N T A
16528 atgtcaagttcgctcttttcaataactgagccttaggttcaagtaggattgattccagtgaccttatattgtttctca
29 M S S S L F L I T E P R S K Y K L G L I P V T L Y C F S
16612 gtttcttttacaggaatgctttcatag 16638
57 V S F T G M L S *
dp1ORF111
28657 gtgactctatcaagaaagctcttgcaattggtgttcaaggttcttgggaaaacttctgcttcttgaagtgacgctgagaaat
1 V T L S R K L L Q L V F K V L G K T S C F L Q V T L R N
28741 tcatcgctgaaaaaacaggtcttcaaatcgctgtctacttaagaaaaattgctcagttcgctgacgctgacaaaacttctgagc
29 S S L K K Q V F K S L S T L R K L L S S L T L T N P L T
28825 ttggttaacattcgctcagttcaactga 28851
57 L V T F V S S T *
dp1ORF112
32207 atgcaaaactgatttaggcaaaactgcttcgacgcagcagccgttgccttatattagatatttgaggaagacaagactcctagc
1 M Q T D L G K Y C F D A A A V A Y I R Y L Q E D K T P R
32291 tatcctggtgacgaaaaagaaaaatccaggattgcaaatgcttatggagtgga 32341

386

29 Y P G D E K K N P G L Q M L M E *
dp1ORF113
17715 atgaaaacagtttaagaagcaatcaaacattcggtgatgaatgggtgacgaaattatcaacgaaaacggccaaatgattcaa
1 M K T V K E A I K Q F G D E W W Y E I I N E N G Q M I Q
17631 gacggaagaatcgaagacatgggcgaatacatggaagaaacggctcgaccaagttaagttcatcaactatggtgacatcgaatct
29 D G R I E D M G E Y M E E T V D Q V K F I N Y G D I E S

17547 caaattatcaaaactatatatcgcataa 17521
57 Q I I K L Y I A *
dp1ORF114
52952 atgctattggcgaagacggggaacagtcctcctgataattgtccattatgccaaaacggattccctcgtattgaaaaactat
1 M L L A K T G K Q S I L I I V H Y A K T D S L V L K N Y
53036 ttcttcaactttacaacctgatacgggaaaagtgaacatgggaccgagccgttcttatgttcaaaagattgttacattta
29 F F N F T T M I R E K L K H G T E A V L M F K R L L H L
53120 tcaataaatatggaagccttgta 53143
57 S I N M E A L *
dp1ORF115
5342 atgagcctccttttttgatatataataacacgaattatcgcgagtttgtaaagccgtttctaaataatttttaactctttt
1 M S L L F L I Y I I Y T N Y R E F V K P F L N N F K S F
5258 aagcatattgagttttgcttcataagtcctcgttcacggcagcctcttgcatgttgagtaaatgaaaggaggttcctcgatatt
29 K H I E F C F I S P V H G S L L H F E Y N E R R F L D I
5174 gttgaaactatagaaggtgaataa 5151
57 V E T I E G E *
dp1ORF116
20662 atgaaattttcaaaactttgctaagcacttactaataacctaagtgtagtgaacaatgaccaagctgaagcttaggcgca
1 M K F S N F A K A L T N E Y L M V V N N D Q A E V L G A
20578 ggaatatcgaaaacttctcaacggttcgaactttgctaattgttagctgaagcgacagttttaaaactcgaaaactcagc
29 G N I E N I L N G S N F A N V V A E A T V L K L E K L S
20494 gaagaggaagctattgagtag 20474
57 E E E A I E *
dp1ORF117
24680 atgataacagcgctgctcgaacattttaaatcgaagtgaatctcgtaagtcactaataagttttgttcaagttatctgctactgtg
1 M I T G C S N I L N R S E S R K S L I V L F K L S A T V
24596 ataaggtctttgacatcgcttgcctcgatatgtcattagtcattggttcattaagaataactcgacaaggaattgttcaag
29 I R S L T S L V P Y M S L V N G S L R I T R Q G I C F K
24512 ccggttggggcggaattcttga 24492
57 P V G A D S *
dp1ORF118
15023 atgatattatctacgtcgacgcaacttgtgaaactattaaatcagaggagcctattgcatgaacaatcagcgaaagcaaatgaa
1 M I L S T S T Q L V K L L N T R S L L H E Q S A K A N E
15107 caaacgaatcgtaacttcgcgaagactatcaacgtgcaagaggtcgaaataaacttccttctgtgttaaaggaccacggcga
29 Q T N R R T S R R L S T C K R S N K L P S C C K G P R R
15191 agaactcgaaaacttga 15208
57 R T R K P *
dp1ORF119
41054 atggagggttaacatccccgattcagtcagtcctactttttcgggcatttcttttagtagacacgacttcagcgggttcgacagat
1 M E V Q H P R F S T S Y F F G H F F S R H D F S G S T D
41138 tttaacaggggaacttcctccaaatcatgtcgcaacattcaagtcaacttcaacaatgcttcggcgcttacggtaccactat
29 F N R E Q L P P N H V E H S S Q L Q Q C F R R L R I H Y
41222 ccaagcatttcacgctga 41239
57 P S I S R *
dp1ORF120
28387 gtgtgaagcgcaagcagaatacatgcgtatgcaattgttcaatcgggtaaatcactgtcaaatcaactaacagcgaggctc
1 V L K R K Q N T C V C N C F N T V N S L S N Q L T A R L
28471 aatacacttacgactacaacatggatgctaagcaacaatcagtcactaagaatggactaaccagctgaaagtgacctta
29 N T L T T T T W M L S N N M Q S L R N G L T Q L K V T L
28555 tcgctgacatttttag 28569
57 S L T F *
dp1ORF121
39222 gtgcagacggatcagtgagttcagtttggaagataataatcaacaatatatgggttattactccgattatgagcaagcagata
1 V Q T D H V S S V W K I I I N N I W V I T P I M S K Q I
39306 gcagggatcgaaactaagtatcgatggttgaccgccttgccaatgttcaagtgggaggtcgaaacgagttccttcaattctttat
29 A G I E L S I D G L T A L P M F K W E V E T S S L I L Y
39390 ttgaatttggttaa 39404
57 L N L V *
dp1ORF122
40402 atgttattctccttatcctacatccgaatcacgttcctgctggattaaacgagattgttccgttctaaatcgccgacttg
1 M L F S L S Y I P N H V H V W I K R V L F R S K S A D L
40318 aatggattgggtaaagatcccggttatcgatgtgaatgaacccttgctgaaggtacataacttcattccctcgaggagaacataga
29 N G L G K D P V I D V N E P L R K V H N F I P C G _ E - H R
40234 aattcggtcacttga 40220
57 N S V T *
dp1ORF123
21327 atgggttcgacttttcgaaggattgaggttttcgaaccggttgagtttttcgagcattctcgacttttcgacccctttctatgct

387

1 M V R L F E G L R F S N R L S P S S I L D F S T P F Y A
21243 cgacttttcgagtggttttgaggttttcgagcaggttcgacttttcgagaaattgagtttttcgacctctaaattaggtcgtgatt
29 R L F E C F E V F E Q V R L F E K L S F S T S K L G S I
21159 attcgaaaagttag 21145
57 I R K V *
dp1ORF124
17891 atggtaaaagttaaagatttgcagtaggaatgaaagttgtaaatgcaaaaggtactgaatttaaagtaactgaccgtcaaggt
1 M V K V K D L Q V G M K V V N A K G T E F K V T D R Q G
17807 cgtaaatgggttaagcctagaacgtcttagtgatggacgtattcggttctatgataacgaatcactaatggacgaaaaagtggag
29 R K W V S L E R L S D G R I R F Y D N E S L M D E K V E
17723 gtagtaaaatga 17712
57 V V K *
dp1ORF125
49916 atgtcctcagccgcttccgtttaaattggaacaagtgaattatatagatgctcctcttttagcttgctgataaggtattcatca
1 M S S A A S V K I G T S E L Y R C S S F S L S I R Y S S
49832 gtttcgccaatttcgaaaaattcgaatccaggaataatgggtcgagaatagtttcgctcgctcggaactcttccatctatcgaaaag
29 V S P I S K N S N P G K W S R I V S S S G T L P Y L E K
49748 tgttcttga 49740
57 C S *
dp1ORF126
16136 atgagctcaagtcagtttctcgaacaatagggtcaagtcagtttatatcaacgaactgtatatcgctcctcttgataggaata
1 M S S T F S R T I G S S P V I S T N C I S S C I G I
16052 aggtctgctacagttgcatgggtgaccctttaaattggagtaactgttccttctactgtttattttaataaggttatctattct
29 R S A Y S C M A D P L I G V T V P S L F I L N K V I I S
15968 atcctctaa 15960
57 I L *
dp1ORF127
13511 atgtaaaatagctttccattccagctcgtgttcttgcgccatttttcagtttcacgatactgaccaactttgcaaaaggtcgt
1 M L N S F P I H R R C S C A I F Q F H D T D Q L C K G R
13427 gaaatagtgctacgattgcaactgtttccattgggttaaattgttctccagccttgcctaccatgggtatccatttcgaaaagta
29 E I V L R L Q L F P L G K C L P S L C L P W Y P F R K V
13343 gttgattga 13335
57 V D *
dp1ORF128
4852 atgacagcagttcaacaagttaagttctacttagaagaagccggcgctcactttctaaaagatggtgagtagcagtaactta
1 M T A V Q Q V K F Y L E E A G A H F L K D V E Y S D N L
4936 gagcaagcaattatgaaagattcttaaatggaatggcgctcatagagatgagcacgatatgaaataaacttcacacgaagta
29 E Q A I M K D I L K W N G A H R D E H D M K I T S Y E V
5020 ttatag 5025
57 L *
dp1ORF129
25133 atgaactttctgctaagcaacttgcgctcactggaagttcaaactaatgtacgcagccaccaatcttacattgaagaattcagta
1 M N F L L S N L R S L K F K L M Y A A T N L T L K N S V
25217 agaaggaagggcggaaggaatgggaacgcattttggaagaactgtcagcttgacgaaatctcagctggagcattgcctg
29 R R K R R T R N G N A F W K N L L S L T K S Q L E H C L
25301 tattag 25306
57 Y *
dp1ORF130
16789 gtgcttgactttattcctttattatcgtataatcataatataaataaaacaagcgtcaaggacgcagaaagaggtcaattatgg
1 V L D F I P L L S Y N H N I N K T S V K D A E R G Q L W
16705 aaacaacactttatttcggttatcttacagcagattggaagacggtcacagaactacactttccactatgaaagcattcctg
29 K Q H F I S V I L Q Q I G K T V T R T T L S T M K A F L
16621 taa 16619
57 *
dp1ORF131
43846 atgctcaaccggctgagaagaaacttggctggcagaagatgctactggtttctggtacgctcgagcaaacggaacttatccaa
1 M L N R L R R N L A G R K M L L V S G T L E Q T E L I Q
43930 aagatgagttcgagtatatcgagaagaaacagtccttggttctactttgacgaccaaggctacatgctcgtcgagaatggttga
44013
29 K M S S S I S K K T S L G S T L T T K A T C S L R N G *
dp1ORF132
15304 gtgactggaaggtcatctaacacatagcctcaagacatttcgttggctttcaggaaaacattcgactagattgtcaatgtat
1 V T G R S S N T H S L K T F R W L S G K H S T R L S M Y
15220 cccacaaaggcttcaagggttttcgagttcttcgccgtggtcctttacagcaagaaggaagtttattcgacctcttgacgttga
15137
29 P T K A S R F S S S S P W S F T A R R K F I R P L A R *
dp1ORF133
8061 atgacttcttcattcatgacaagttttcgagttttctgcttgcttgctcaggaatagttttcccgccggtcaaaatgtatagatt
1 M T S S F M T S F R V S A C L S G I V F P A A K M Y R L
7977 tcgtatttttcttctgtagcagaacttgaaatccatttgcattccaccatttccgacctatcgcggcgaaataa 7900
29 S Y F S F L I A E L E S I C I P T I S A L S A A K *
dp1ORF134
498 atgacttcaatgtacttaggttccatcaattcatacaagtcattcaaaataatgttcatgcaatcttcgtggaagtcaccgtg
1 M T S M Y L G S I N S Y K S P K I M F M Q S S W K S P W
414 ttacggaaactgaataagtaaatattcaatgatttagattcaaccatcttttcgttggatgtaa 349

389

29 S K S K G I V F E I L I F M S S R F P *
dp1ORF150
15185 gtggctctttacagcaagaaggatttattcgacctcttgacgcttgatagctcttcgcaagttcgacgattcgtttgtcat
1 V V L Y S K K E V Y S T S C T L I V F A K F D D S F V H
15101 ttgctttcgctgattgttcagcaataggctcctcgattttaaataagtttcacaagttcgctcgacgtag 15033
29 L L S L I V H A I G S S Y L I V S Q V A S T *
dp1ORF151
28027 atgattatatcaacgcaggaggagattgctagctacattcaagcacttcttcaaacgctcttcaataccttggaaccaactcttt
1 M I I S T Q G R L L A T F K H F L Q T L F N T L D Q L F
28111 tccctaattgctcaacaaacaggacagacatttcatggctcaagggcgcaataatttgcagtag 28176
29 S L M L N K Q G T F H G S R V Q I I C Q *
dp1ORF152
42235 atgtgcataaaggacttatcgacaaaggaggtactattgcagctacttctgaaggatttagaccgaaagtttcaatgtatcttc
1 M C I K D L S T K R L L L Q Y F L K D L D R K F Q C I F
42319 aggtcttcaataactcatatggaatgccatttcatgtatatacactgacggaagacttgtggtga 42384
29 R L S I T H M E M P F Y V Y T L T E D L W *
dp1ORF153
22307 atgggtggacaaagggtcaccttttgcgaacttctcgatctcgatagcagacgggtccattcggttcaggaaaaacagtatcgat
1 M V D K G L T F S N F R Y R H S R R F H S F R K N S I D
22391 ggctctttcattttcccttttggccatgacggaattcaacggacaaaactttgccatctgtggtga 22456
29 G S F I F P L G H D G I Q R T K L C H L W *
dp1ORF154
18446 gtgacaataggctttaagaactgcaaaaaaacctggggcgctgcacgcgcaaacctggagctccttaacagtcattccaaggctg
1 V T I G F K N C K K T W G V C T R N L E L L N S H P R L
18530 aggtttcttacaacaatcctaattccttcaaaatagctcttgcgggtcaatagtgctaa 18592
29 R F L T N N P N S F K I A L V R V N S A *
dp1ORF155
13512 atgaatacgcacctgagcaacttacaatgggacatgggtgcaaaatctaatttcttctcaacgtttcattcaactcacgcca
1 M N T T L S N L Q W D M V Q N L I S F F N V S F N S R Q
13596 ttgaagctcaagcaattttctggcatatgggagcctatgatattagctccttatgcaaatgtga 13658
29 L K L K Q F S I W E P M I L V L M Q I *
dp1ORF156
18777 atgctagtagtctccatttctgttggcttctgtttagctctgttcagttcagctgcttctcgcatgcaatagtttcgagaat
1 M L V S P F L L V L L F S S V Q F S C P S R C N S F E N
18861 atgcctgttcataggctcacaatattccgccaagatttgcagttatggtggcgtaattaa 18923
29 M P V H R L T I F R Q R F A S Y G G V N *
dp1ORF157
13281 gtgcttctgctggacttgagaagaatttggtatcttttcgagccaatccataagggttctcgataccgtcacgattgattgttct
1 V L A G L E K K L V S F S S Q S I R F S I P S R L I V S
13197 gttactgctttcttgaagcgttttttaagctgtgcatattagaccctttcattttctataa 13135
29 V T A F L K R F L K S V I L D P F H F L *
dp1ORF158
40727 gtgaacgcggttattagggtcaaacgaagcccaacggacattgtcttctgtcccgctcactattgtgaggaaacagtcacttctcc
1 V N A V I R V K R S P N G H C L C P V T I V R N S H F S
40643 acttgcgagcgttacctcttcgcccagctgtcgttagctggtgactgctatgaacattga 40581
29 T C E R Y L F A G R V V V W V T A M N T *
dp1ORF159
30371 atgatttggctcgcttaccacagcagcttctcctttgagtttctgtcgagcattccctgtacgggtctgtccaaatagcagc
1 M I W S A L T Q A A S P L S F C R A P P V R S V Q I A C
30287 gtcttttgcgtatttcttcattcttagtagcagcagcttgcgagactgttatgacagcagcttga 30225
29 V F A Y S S I L V A A T S Q T V M T A T *
dp1ORF160
41324 atgggttacagacacgcgaggaacaaatcgaacgtccaagcgtatctatcaatgttatagaataactatggaccgtctatcaa
1 M G Y R H A R K T I E R P R R I Y Q C Y R I L W T V Y Q
41408 tttctccgttcaacgtactcgtcaaaatcctgcaattatccaagctcttcgaaatgctaa 41467
29 F L R S T Y S S K S C N Y P S S S K C *
dp1ORF161
52175 atgcaaaaagggtttaaatgcttatctcgacatgacattgaaagcattgcattcgagactatttcaaaatggttgcaacgttca
1 M Q K G L N A Y L D M T L K A L H S R L F Q N V W Q R S
52259 aatcaaaccaaggggccaagttttcaacttaccttacaagactcttcaagaatagaatag 52318
29 N Q T K G P S F Q L T L Q D S S R I E *
dp1ORF162
13020 atgacagaagttcggttaaatagcccgcaaaagggtgagagtagttatgggtcggaatattgaatttctcgaatattttaaaagg
1 M T E V A V N S P Q K V R V V M V G N I E F L E Y L K R
13104 aagtacggaacagaaacttccatcagttatattatagaaaatgaaagggttctaataatga 13163
29 K Y G T E T S I S Y I I E N E R G L I *
dp1ORF163
40224 gtgaccgaatttctatgttctccgcaggggaatgaagttatgtaccttacgcaagggttcattcacatcgataacgggattcttta
1 V T E F L C S P Q G M K L C T L R K G S F T S I T G S L
40308 cccaatccattcaagtcggcgatttagaaggaacaaatcctggttaattccagacatga 40367
29 P N P F K S A D L E R N N T R L I Q T *
dp1ORF164
6696 atgtactcttggagaacttctgctctaaatgttccagcttcccccattgcaattaggttagaattctgcgttatctataatagac
1 M Y S W R T S C L N V P A S P I A I R L E S A L S I I D
6612 tcaccgattctttcgaataacatttttgaatacattccaccaaccccgctgggttataa 6553
29 S P I L S K Y I F R I H P P T P L G L *

dp1ORF165
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1 M S E S W S I P T T D G L Y L D I M L S K I A G V R F F
50420 cctccaatcataaaggcggtgactaccacaagggaattttcagcctcagtcattgcttga 50361
29 P P I I K G V T T T R E F S A S V I A *

dp1ORF166
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1 V V M L F N D S I F S R L A R F T V P A V S I V F I N V
23435 gtgcgtgttgcagggtcgagtgtaaatctattctcagccaagagttcagcgtgaaatga 23376
29 V R V A R V E C K S I L S Q E F S V K *

dp1ORF167
1008 atgcttattcgggtggagcttcttacctcgtatattggctcagcagacgatgcggctggaggtgcttaccctgattgcactc
1 M L I R L E L L T S Y M V L T Q T M R L E V L T L I A L
1092 ctgagttctataattcaatgcaaatgcaatggaatggaactggaggcaaggtaa 1148
29 L S S I I Q C Q M Q W N M E L E A R *

dp1ORF168
54345 atgagactttttccagggttatattcttcacattgttcagttcctggagtcaggtattgttctttaaattcatagagttcgaaag
1 M R L F P G Y I L H I V Q F L E S S I V L E I H R V R K
54261 ttgcaagggtcatagccgcacacataggaacacaggaattaaactaa 54205
29 F A K G H R P H T Y R Q H Q E E L N *

dp1ORF169
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1 M N T A S R R V S M L V I R K N S S W P P S K S S A R L
45870 gaaactccgtcaatcactaatcttccatctttagtgactcgaactcctaaatatga 45814
29 E T P S I T N F P S L V T R L P K I *

dp1ORF170
27600 atgatgattgttcttctgtgctcctgccgttggtagcagcagcaagttgcttaccaaaagagccgatttcacgaggttcgggaa
1 M M I V L V L L P F V E Q Q Q V A Y Q K S R F H E V R E
27516 caccaccacgacacgacgtggatttctaaatttcagtcctccgggtggcgacttag 27460
29 H H H R H D L D F L N F Q S R L A T *

dp1ORF171
47678 atgtcatctttcttcatgtactcttttagagcatcacgaagacttttgacttgtttctccatgtcgccttggtagcattta
1 M S F S F M Y S F R A S R R L L T C F S M S P L V A F N
47594 tcaccggtcttctcaattgcagcgatgaactgttttctcatcttcaaatcttcaat 47538
29 S P A S S I A A M N C F S S S N F I *

dp1ORF172
10462 atgtttcgaacattttctacccattattagaagcagcatcaatttcaataggagagccaagtccttctgttcacatccttcg
1 M F R T F S T P L L E A A S I S I G E P S P L F T S F A
10378 aaaattcgagcagtagtggtttaccagttccagcgcaccacagaatagatag 10325
29 K I R A V V V L P V P A P P Q N R *

dp1ORF173
32160 atgacattagacatttcttctgtctgtacgaaagggtttcagcttgagtcacttcacgtacattgcactgaagattgtcataag
1 M T L D I S F V C T K G F S L S H F T V H C T E D C H K
32076 ttgctcatctgtcatatactcgcgacttcagcgtgaagtaggctctaccattga 32023
29 L L I C H I L A D F S V S R L Y H *

dp1ORF174
29766 atgtcccatcagcccttttcatgaagattgtcgaaccagcgttcgacttttcatcagtttcaagctgttcttgcattatattgt
1 M S H Q P F S L R L S N Q R S T F H Q F Q A V L A Y I G
29682 cataatgaattgcgccatttgttccagtagtctgcgtcaccttttagactga 29629
29 H N R I A P F V S S S L R H L L D *

dp1ORF175
15648 atgcgcgtgatgtcatggcagataggcgaggataaagagtgctgaatagaacgccgagagcttacgagagcgccaaatacaag
1 M R V M S W Q I G E D K E C R I E R R R A Y E S A K Y K
15564 ggccaggtactacgggtggtccttctgcttacctgtaacaaataaaccattga 15511
29 G D G T T V V L L L T C N Q I N H *

dp1ORF176
43031 gtgataaagacggtaacgttgaattttctagttcgtcttgaatgacgtcattttgggtgattgattgctactgtcgtttggct
1 V I K T V T L N F S S S V L N D V I L V I D C Y C R L V
42947 aatcccgctcgactgctgtttaaagagtgtaagagttgtagagatatcctctaa 42894
29 N P V D L L F K S A K S C R D I L *

dp1ORF177
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1 M N L N S S R L L K L L G K K Q V E Y F G G N V N L V I
19853 ttctcgcgactaatttttaggtgcttttgtattaatcagcgtgatatgcgcttga 19800
29 F S R L I L G A F V L I S V I C A *

dp1ORF178
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1 M T T V D Q F K R Q L R K S L G S I F P S S V - S L - N - L S
11840 caattagtaaccttttagcgaattgttagcacttgcctcccatattaagtcataa 11787
29 Q L V T F S E L L A L A S H I K S *

dp1ORF179
56058 atgggttaggggttatcttacctcgttgattgttcttatgcaaaacctaccacaatcgcttgcgtgggttcaggagttgcatt
1 M G R V I P Y L V D L L Y A K P T T I A C R G F R S C I
56142 ttggataagtcaaaaagcaagtgcttcttatattcgacaagctctcgaataa 56192

391

29 L D K S K S K C L Y I R Q A L E *
dp1ORF180
41176 atgttcgacatgatttggaggaagtgttccctgttaaaatctgtcgaaccgctgaagtcgtgtctactaaagaaatgcccga
1 M F D M I W R K L F P V K I C R T A E V V S T K E M P E
41092 aaagtaggcgtactgaatcggggatgttgaacctccatccgtttgaatag 41042
29 K V G R T E S G M L N L H P F E *
dp1ORF181
13126 atggaagtgttctgttccgtacttcccttttaaatattcgagaaattcaatattcccgaccataactactctcaccttttgcggg
1 M E V S V P Y F L F K Y S R N S I F P T I T T L T F C G
13042 ctatttaccgcaacttctgtcataggctgtcctcctttgtcttatactgtaa 12992
29 L F T A T S V I G C P P L L I L *
dp1ORF182
45369 gtgcttccccatgtttcaataaattaggggtcgacctgcctagctttcgaacgtgctataacgatttcaatcatagcgaagaaa
1 V L A H V S I N R V R P R L A F E R A I T I S I I A K K
45285 ggtgagaagcttcaatcaattccattgcgggtgcaatatcttcttctga 45235
29 G E K L Q S I P L R C Q Y L L P *
dp1ORF183
13896 gtgattccagcttttgggttttcttcagcctcttcaacttttcttcttaggcgaggtttctttaggtgaactcttaggt
1 V I P A F G F S S A S S T F S S L G A G F L R V E L L G
13812 ttttcttcaactacttcttcaacctcagcctcttcttcaactggacctga 13762
29 F S S T T S S T S A S C S T G P *
dp1ORF184
53330 gtgaacttgcgctcaaccacgtcaaactattgttcttctgcgaggtctaaaattagagttccaagaagttcgctcttttctgga
1 V N L P S T T S N I W S S S R S K I R V P R S S L F S G
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29 K S S R V A L S S G R S G R N S *

dp1ORF185
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1 M K F E M F E M K I Y L L L' D T L E M A K K L S T T S I
22606 tatttggaggaaaagatgagtcgagtcagacattatcacgggggttaa 22653
29 Y L E E K M S R V K T L Y R G *
dp1ORF186
21272 atgctcgaaaaactcaaccggttcgaaaaactcaatccttcgaaaagtcgaaccattcgaaaagttcaaaagttcgaaaaactc
1 M L E K L N R F E N L N P S K S R T I R K V Q K F E K L
21356 aaccattcgagagtaggaattaaggacataaccaggtcgaaccttttag 21403
29 N H S R V G I K D I P V Q P F *
dp1ORF187
34415 atggctctgttcaatctcttctactatcattcaagcagctgttcaaaattatcactgcttattcaatggctgttcttcaggcag
1 M V L F N L F L L S F K Q L F K L S L L Y S M V L F R H
34499 ttctcagcttattcaagcaggtcttcaaaatttgcagctctcataa 34546
29 F L R L F K Q V F K F C Q L S *
dp1ORF188
35609 atgttcgtaaaagcagcgggttcgacctgagtggaactgttcaatacaggaagtgcacaacctaaccaacctcagtcacaatcta
1 M F V K Q P V R L E W T C S I Q E V T T L T N L S H N L
35693 aaaacaatcaaggcgagcaaacgttgcacattggaacaatcgtag 35740
29 K T I K A S K P L S T L E Q S *
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42587 atgcaaacgcagtatcaaccgtctctgaaactcttcatgaccagacttgatgtgctggaaccgtcgagaacttcgagctgacg
1 M Q T Q Y Q P S L K L F M T Q T C M L R T V E N F E L T
42671 agcaaaaacttcgcaaaactcggtacgcaatcgaagatgaaattctag 42718
29 S K N F A K L V T Q S K M K F *
dp1ORF190
39786 atgtattcactcaaaagttgttcagtggtgctcaatcatattaaaatcgaaacttgtaatatcttactccttttagtgaagcag
1 M Y S L K V V Q C G S I I L K S N L V I S L L L L V K Q
39870 aggaagaccttaaatatcgaattgactcaaaagccgatcaaaagctaa 39917
29 R K T L N I E L T Q K P I K S *
dp1ORF191
40996 atgtccattgttccggaacttgatttaggtaagtaccttgtaagtccagtgacggcgtaaggatagctagtagtattggttc
1 M S I V P E L D L G K Y L A K S S D G V K D T L V V W F
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29 L P K S I Q S L P K T R Y Q T *
dp1ORF192
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1 M V D V E C F F E M K F R V F S I P Y G M F S E C F N K
2836 acggaatggagtattcttgcacccgtcacgttctgctcctcgcttaa 2789
29 T E W S I L Q P V T F C V L A *
dp1ORF193
42456 atgatttcagctcaaaatgaaatcgaatgagacattgtctaaatttaaccaagaattatctacattcgatttcacacaaagt
1 M I S A Q I K Y E M R H L C L N L T K N Y L H S L S P Q V
42372 ttccgtcagtgatatacatagaatggcatttccataggtattga 42325
29 F R Q C I Y I E W H F H M S Y *
dp1ORF194
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1 M N P C V R Y I T S F P A E N I E I R S L D T L M V E L

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40200 ccgctcgttcttaccgataaattagaccttcattagaagagctcatgtaa 40153
29 P S F L P I I R P S L E E L M *
dp1ORF195
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1 M F T I V V L T S F F S A P C P I V N S A T I W R D F V
42500 aggttcaacatagttctcacctcctttctaaaaaatattataacatga 42453
29 R F N I V L T S F L K N I I T *
dp1ORF196
11273 atggtagatttaacaagtcctcgtccaatcatgtcactcctccttgcctcatcaaaagaagtttggtttcaattatcggttttagc
1 M V D L T S P C P I M S L L L A H Q K K F G F N Y R F S
11189 attaggctcccatTTTaaactccagcaagttcattcatttcttcttag 11142
29 I R L P F N N S S K F I H F F *
dp1ORF197
7484 atgaaaagattataggtatccaatttcaagccttgaaaaaattaaacggctctggagttaaagcgctcaacccaaacttcatcg
1 M K R L Y G I Q F Q A L K K L N G L E L K A S T Q T S S
7568 atgcagggtatgaagtttcttacaagaagcgctgaactagatga 7612
29 M Q G M K F L T R S V E L D *

dp1ORF198
24119 atgccgctcaacaaattgacgtccagttttattcaatgcctcagttcacctatacagttgaccctagaacccttccagcttgc
1 M P L N K L T S S F I Q C L S S P I Q L T L E T L P A C
24203 tttctgttgacattgtttatcaggacgagcgtacaaaaggaaatga 24247
29 F L L T L F I R T S V Q K E *
dp1ORF199
15742 gtggcctcctgaattaggtgtacttttctcccaactgcttagcaactgccttctctgttttagcactagctctgcgcgtggga
1 V A P E L G C T F P P N C L A T A F S C L A L A L R V G
15658 attggtttgtatgcgcgtgatgtcattggcagataggcaggataa 15614
29 I G L Y A R D V M A D R R G *
dp1ORF200
47843 atgacaggcttgattcgataagccctgaaagtttttcacacatttcttccgtctcgtccttgcgaactaatttttcgataatt
1 M T G L Y S I S P E S F S H I S S V S A S S T N F S I I
47759 tctttcaagcgttcttctcgtccatagttgagcgtctcgtcgttag 47715
29 S F K R S S S I V E R S V V *
dp1ORF201
38569 atgggcttcacaagttccttctttaaataaagggtcaatatctttggactcgaactatttggacctataccgattcaactaccga
1 M G F T S S F F N Q R S I S L D S N Y L D L Y R F N Y R
38653 aacgggctatcaaaaaactacattccaaaagacgggaatga 38694
29 N G L S K N L H S K R R E *
dp1ORF202
44483 gtggggcggtttattttttataaaaaattttttacaaaatgcttgacaacattcactcattatcgataatacaattataaaaaata
1 V G R L F F I K I F Y K M L D N I H S L S Y N T I I K I
44567 aataaaagccgaaaggcgaggagacattatgtcaaaaattaa 44608
29 N K A E R R G G H Y V K N *
dp1ORF203
22781 gtgattaggattggcgggtttacaagagaaccacattttcgaacctgttacggaacagcgccctgtcgttgggtgacaaacga
1 V I R I G R V T R E P H F R T C Y G T A P C R L V D K R
22697 ttcaggcatcagtgccacctcaccagaaagatacctgctaa 22656
29 F R H Q C H L I T E D T C *
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1471 atgaccacggttcgagtcagggtggtgttgacttttatcacgtcaagaaaatcgaggtacattcattgacagacttgacc
1 M T T V R V K G W L L T F I T S R K S Q V H S L T D L T
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29 T L F F F K G M N Q S L *
dp1ORF205
8524 gtgacactgatgaatggttctcagtttgggtatgctactcgtgacgcagatatcttctacgaccaaagaattgcccaatttagaa
1 V T L M N G S Q F G M L L V T Q I S S T T K E L P N L E
8608 ttcaggaagcaacctgctatcaagttcaatttcgtag 8646
29 F R K S N L L S S S I S *
dp1ORF206
19855 atgaccaagttcacgttcccacaaaatattcgacctgcttcttcccaacagcttgagaagctctcgaactgttttaggttcac
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19939 aaattgttcaacttgagcaagtgcgatattattctttag 19977
29 K L F N L S K C D I I L *
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27502 gtgtcgtgtgtgtgttcccgaacctcgtgaaatcggtcttcttggtaagcaacttgctgctgctcaacaaacggcaggagcac
1 V S V V V F P N L V K S A L L V S N L L L L N K R Q E H
27586 aagaacaatcatcattctttaaataataggaggaaactaa 27624
29 K N N H H S L N N R R N *
dp1ORF208
47279 atgtttggtatgaagcaaaagacttcgtgaagaaaataacattcacttcccgtttgttcttctcctgaacctagaacagaccttg
1 M F G M K Q K T S L K K I T F T S R L F F L N L E Q T L
47363 accatcgtgttctcgtattctgggatgacgaaggcgtga 47401
29 T I V V L D S G M T K A *
dp1ORF209
29784 atgttaagaatcaagttcgtagaccattgaaacgcctcctactaaaatcaaggtacttcgaaactcttgggtcagtgatggat

393

1 M L R I K F V E P L K P L L L K S R Y F E T L G S V M D
29868 atggaggaaaagataaaagcgaatgaagtcgtag 29906
29 M E E R K R I K R M K S *
dp1ORF210
53077 atgtttcaacttttccgctatcatggttgtaaagttgaagaaatagttttcaatcagagggaatccgttttggcataatggac
1 M F Q L F P Y H G C K V E E I V F Q Y E G I R F G I M D
52993 aattatcaggatggactgtttcccgctcttcgccaatag 52955
29 N Y Q D G L F P R L R Q *

dp1ORF211
20959 gtgctcgacttttatgtcgcccctaattttgtttttacttacggactatgggattttaggtattttcagggcgctttttat
1 V L D F Y V A P N F C F Y L R T M G F V G I F R A L F Y
20875 ttacttattaagtccttttctatattagattgtttataa 20837
29 L L I K S F S I L D C L *
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52983 atggactgtttcccgctcttcgccaatagcattgcaattgatatagcgtcgacgaccgtcaacgtctgcttcgtggactacgaa
1 M D C F P V F A N S I A I D I A S T T V N V C F V D Y E
52899 ataaccatgtcttcgcttcgggtcatcacaatag 52861
29 I I H V F A F R V I I Q *
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30291 atgcgtctttgctattcttccatcttagtagcagcgacttcgcagactgttatgacagcgacttgaaactgtttcgataccg
1 M R L C V F F H L S S S D F A D C Y D S D L K L V S I P
30207 ttcacagttactaacaattcttcaggcttcatactaa 30169
29 F T V T N K F F R L P Y *
dp1ORF214
24273 atgatgccaaagtgtttttcagtgctcattcttttgtacgctcgtctgataaacaatgtcaacagaaagcaagctggaagg
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24189 gtttctagggtaactgtataggtgaactgaggcattga 24151
29 V S R V N C I G E L R H *
dp1ORF215
35822 atgttaccaaaccctgatagagtttcttacttctattatacaatcctctcgacagtttgtcaacgtcgtcattgtttcgaact
1 M L P N P D R V S L L L L Y N P L D S L S T S S L F R T
35738 acgattgttccaatgttgacaacgggtttgctcgcctga 35700
29 T I V P M L T T V C S P *
dp1ORF216
32849 atggcctcggagctcgcggccacatctctccagatagcggcagcagggtcaagtaccctggcatagcgtccatgatttcattt
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32765 acctggaaccggctgaagctagattttccatacctga 32727
29 T W K P A E A R F S I P *
dp1ORF217
23443 atgaatactatgcttacagctgggacagtaaaagcgagccaaacgggagagaagatagagtcattaaagagcatgaccactgcatgg
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23527 ataggaacagatagcctgtctcactgacgctctaa 23562
29 I G T D M P V S L T L *
dp1ORF218
22029 atggaatgcttccggaagaggttcgatataactacaaattgagcgcgagaaaattacattgctccgggccaatggcgacc
1 M E C F R K R P D I D Y K L S A R K L H C S G P K W A T
22113 aggaaattgaaggcgaggttaaagataacttcgtag 22148
29 R K L K A R L K I T S *
dp1ORF219
51388 atgattttatgctcgactttttcagttctccatttcttcgaaacgcttcagggtgacgccttgcttaactacttcgctagat
1 M I L C S T F S V L P F L R N A S G L T P C L T T S L D
51304 gttccaaaattccttttcagccactgggttccatag 51269
29 V P K F L F S H W F P *
dp1ORF220
6334 gtgaagttttcttcggtgacggttgatacaatttcttcaagagtaagctgttaagggtggcaagtgaattctttctcgaaact
1 V K F S S V T V D T I S F K S K L L R W Q V N S F F E T
6250 ttcttgccagcagatgcgtacatgatgtcttcataa 6215
29 F L P A D A Y M M S *
dp1ORF221
43507 atgactgctcaagttctatgtactatgctctccgctcagccggagcttcaagtgtggtgggagtcataactgagtagatgc
1 M T A Q V L C T M L S A Q P E L Q V L D G Q S I L S T C
43591 acgcatggcttattgaaaacggttatgaactaa 43623
29 T H G L L K T V M N *
dp1ORF222
13212 gtgacggtatcgagaaccttatggattggctcgaaaatgataccaatttcttctcaagtcagcaagcactcgataccatggaa
1 V T V S R T L W I G S K M I P I S S Q V Q Q A L D T M E
13296 gctatgaagggtgacttgcgagcactcattaa 13328
29 A M K V D L S S T H *
dp1ORF223
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1 M W W Y L L D M F E M S T T S T V K S L T F T T R K M S
14139 acgagcctgacgatgacagcgacattctttag 14171
29 T S L T M T A T F L *

dp1ORF224
13621 atgccagaaaattgcttgagcttcaactggcgtgagttgaatgaaacgttgaagaaggaaattagattttgaccatgtcccat
1 M P E N C L S F N W R E L N E T L K K E I R F C T M S H
13537 tgtaagttgctcagggctgattccatagctaa 13505
29 C K L L R V V F I C *
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32991 gtgagcaacgggtgcgacgtatttcatcgccctctgccatgtcgctagtttctgcgttcgtatcagctgctgctcgagcaatac
1 V S N G C D V F H R L C H V A S F C V R I S C C S S K Y
32907 gtcagccacgtgacccgctggtttgctctaa 32875
29 V S H V T R L V C L *
dp1ORF226
25191 gtggctgcgtacattagtttgaacttcagtgagcgcaagttgcttagcagaaagttcatcgctaggaattggatagtggtgttc
1 V A A Y I S L N F S E R K L L S R K F I A R N W I V V F
25107 gatagtcattgtcgtgaagtgtttgataactga 25075
29 D S H C R K C L I T *
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23115 atgactcaattagatggtagcgttatgacgttttcgagaatccataaaggccgaaggtgttgcattatagataccaaagtcgc
1 M T Q L D G S A Y D V S R I H K G R R L L H Y R Y Q S R
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29 L L R I N G R I L Y *
dp1ORF228
10450 atgttcgaaacattattgaagattctagatacaagtcctatggacagcaggttcaaagtttacatcattgacgaggttcattatgc
1 M F E T L L K I L D T S L W T A S S K F T S L T R F I C
10534 tttcaaccggagcatttaagtcgctgttga 10563
29 F Q P E H L M R C *
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27634 atgtgcgagttaagaaaactgattttaatcaaacctcgaagcattgtcgcaattcctgaccactacgttgcgttggtgctc
1 M C E L R K L I L I K P L E A L S Q F L T T T L L W L L
27718 aaattccagctaccgagcaactcaagtag 27747
29 K F Q L P Q Q L K *
dp1ORF230
50723 gtgacgaaaaatccggcacttgaactatctgctgttaaaaaccgatatggcgaagaccgaaaaatcatcgaatatatgtggg
1 V T K N P A Y L N Y L L K T D M A K T E K S S N I C G
50807 acgttgaaactggaacctatactcttatag 50836
29 T L K L E P I L L *
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31071 atgcgcgtgtcattgcgtttcacatcttcagttccctccgaggtcacggcttcgagttctgctgtttctgcgctatctaogaca
1 M R V S L R F T S S V P S E V T A S S S A V S A V S T T
30987 aagttagctccgagcacttttggcaactga 30958
29 K L A P P T F G N *
dp1ORF232
29385 atgtcaattccattagctcttgcataattcaacgagctcaggaacgggttttagccgcatactcttcgcgcatttgttcaacttcg
1 M S I P L A L A N S T S S G T V L A A Y S S R I C S T S
29301 tcaatttcttcaactgattcaattgtttga 29272
29 S I S S T D S I V *
dp1ORF233
52892 atgtcttcgcttccgggtcatcatacaatagagtgaacaattgcgctgtcaccgtggtcagcagtggtgaaaaactcgttatta
1 M S S P S G S S Y N R V T I A L S P W S A S V K N S L L
52808 gaccctgagctaaatgttctctgattttga 52779
29 D P E L N V P D F *
dp1ORF234
36253 atgcttacgagtagacgagcactcaactgttcgaaagggttataagtttcaaccgctttgggaggcgatagcttacctaaccag
1 M L T S T A T Q L F E R F I S F N P L W E A I A Y L T Q
36337 gaagacctactcgacaatttagagtag 36363
29 E D L L D N L E *
dp1ORF235
32768 atgaaatcatggacgctatgccaggggtacttgacctggctgccgtatctggagagatgtggccgcgagctccgaggccatgg
1 M K S W T L C Q G Y L T W L P Y L B E M W P R A P R P W
32852 ctagtccacttcgagcctttggattag 32878
29 L V H P E P L D *
dp1ORF236
37528 atgttcgctcgttttagatttagcaatatatcgagggttcattgtggcgtgtagtaaaccacgaaacatcaatgagatattcact
1 M F V A F R F S N I S R L H V A C S K P R N I N E I F T
37444 tccattgttgatagaagcaaacgttaa 37418
29 S I V D R S K R *
dp1ORF237
1678 gtgagagtcaggtaaggaatcttgacatattctcagccgtagttctaaatccaaatagaactcgcttggtgtcgaactgcattt
1 V R V Q V R N L D I F S A V V L N P N R T R L V - S T A F
1594 gctaaagcgattgggttcattccctga 1568
29 A K A I G S F P *
dp1ORF238
1301 atgcctttttgcggtcgatataagttgcgcaagttccacaactttcagcgtcacttttcataacatgaacgagtcagaaataag
1 M P F C G R Y K L R K F H N F Q R H F H N M N E S R N K

395

1217 gaacatctaaatcaattccccatttaa 1191
29 E H L N Q F P I *
dp1ORF239
26521 atggtgaagtatttccatcgagaatgtcctttcgaccatcctaaggaatgtgctaccaaactgtatggtacgaaaactcac
1 M V K Y F L S K N V L S T I L M E C A T K L Y G T K T H
26605 tcgaagaaatcgctgatgagttga 26628
29 S K K S L M S *
dp1ORF240
41893 atgtttggaataagcgtgaaacagagtttacatggcgaagtaacaaatcgaggacaacccctacgggaactcgaggtgaatggg
1 M F G I S V K Q S L H G E V T N T R T T L R E L E V N G
41977 gactatttcaaaatttctggttag 42000
29 D Y F K I S G *
dp1ORF241
47020 gtgtctttccttaatatggagatagtttctattctatttaagcaggatctgaaaagggtaccaatttttagatttcataggctt
1 V S F L N M E I V F I L F K Q D I E K V T N F R F H R L
46936 accatctacgatataatctgctaa 46913
29 T I Y D I I C *
dp1ORF242
41338 gtgtctgtaacccatgctcttaccgtagcggagccattaaagttcatcatacccaatttgcgcgcttttcggtgatagcttgg
1 V S V T H A L T V A E P L K F I I P N L P P F S L I A W
41254 tttttacctacgagctcagcgtga 41231
29 F L P T S S A *
dp1ORF243
51306 atgttccaaaattccttttcagccactgggtttccatagaacccctccatcgtttcgacctaatatcattcgagacgaattcagtta
1 M F Q N S F S A T G F H R T L H R F D L I H S R R I Q L
51222 gtcctgaagtgttagccgcaagtga 51199
29 V L K C S R K *
dp1ORF244
27083 gtgaggtacaaaaatgttgaccgtcgccgtcaatgaaaatttttagcatcgagttcttttcgaagttttcgaataatttcttcac
1 V R Y K M L T V A V N E N F S I E F F R S F R N N F L H
26999 ctgtttgatagttggttcattctag 26976
29 L F D S W F I *
dp1ORF245
6278 gtggcaagtgaattctttcttcgaaactttcttgcagcagatgcgtacatgatgtcttcataactgctagtagaagttttaat
1 V A S E F F L R N F L A S R C V H D V F I T A S R S F N
6194 tcgaagtcggtctttcagaataaa 6171
29 S K S V F Q E *
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2831 atggagtatcttgcaacccgtcacgttctgcgtcctcgcttaataagacaaaaagtcctttgaacggctgcctcagttattgtcca
1 M E Y L A T R H V L R P R L I D Q K V F E R L P Q Y C P
2747 aggttacaatttcatcggccttaa 2724
29 R L Q F H P A *
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29641 gtgacgcagactactggaacaaatggcgcaattctattatgaccaatataagcaagaacagcttgaaactgatgaaaagtcga
1 V T Q T T G N K W R N S I M T N I S K N S L K L M K S R
29725 acgctggttcgacaatcttaa 29745
29 T L V R Q S *
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53644 acatttcaggaaacgctctaa 53664
29 T F Q E T L *
dp1ORF249
2012 gtggatgcgactatcattgcaactggtgtgactcagcctttacctggaacggtactactgagccggaatatatcacaggcaaag
1 V D A T I I A T G V T Q P L P G T V L L S R N I S Q A K
2096 aagctgctagtcgaatcttga 2116
29 K L L V E S *
dp1ORF250
23837 atgggcaaacatggaagattgacgaagactcagtcgactataaacctactcgagaaattcgaactatattcgacaacttatca
1 M G K H G R L T K T Q S T I N L L E K F E T I F D N L S
23921 aaaagcaatcacgctttatga 23941
29 K S N H A L *
dp1ORF251
39205 atggaataattagttcttaccgtctgcgctgggttcccggtatcccttgagctccgtcattcccttccatttcgtccatgt
1 M E I I S L T V C A W L P G Y P L S S V I P L P F R P C
39121 ataggctgcaggggtcttttga 39101
29 I G C R V F *
dp1ORF252
54771 gtgttgataggtcgaaactaattttgcatattttctatatttcaaaagtgttttgagatatcgttataaaatgctcgacaa
1 V L Y R S K L I L H I F Y I S K V L L R Y R Y Q N A R Q
54687 tactttcgctgttctcttag 54667
29 Y F R L F L *
dp1ORF253
56255 atggttcgctctataatagaaccgatgttgctagacaaagcatttgcaatcttcgagtcctaattttattcgagagcttgctgaat

396

1 M V A S I I E P M L L D K A F A I F E S N L F E S L S N
56171 ataaagacacttgcttttga 56151
29 I K T L A F *
dp1ORF254
48479 atgaacctttcgcttaggttcaatctttttcgaacattttcatatttaacaaaactttcagctaaaaatcgacaaagttcaatg
1 M N L S L R F N L F R T F S Y L T K L S A K N R Q S S M
48395 ttcgactcaatgttttaataa 48375
29 F D S M F K *
dp1ORF255
9572 atgctttggtcttctcgacgaatgactctactacattccctgcagggtttcgagcagtacgggtcaatgatgcaccgttttctg
1 M L W S S R R M T L L H S L Q G F E Q Y G S M M H R F R
9488 caaggtagtcaccttttctaa 9468
29 Q G S H L F *
dp1ORF256
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1 M T F Q S L M R P L K L D T T I H G F T N F E T K Q L K
15373 cacttgaagaaatttttag 15390
29 H L K K F *
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28216 gtgaacgtgctggatttagcaaaacagctactgagatggcatttctccgtgagtcctatgcgacttggtgaaaaagacgtcaaa
1 V N V L D L A N K L L R W H S S V S L C D L V K K T V K
28300 acttgcaaatgctattga 28317
29 T C K C Y *
dp1ORF258
44023 atggaaattggtattggttcgacctgacggatacatggctacgtcatggaaacggattggcgagtcaggtactacttcaatc
1 M E I G I G S T V T D T W L R H G N G L A S H G T T S I
44107 gcgatgggttcaatggtaa 44124
29 A M V Q W *
dp1ORF259
4298 atgactcgactacgaagcataaagacaagtggatggaaagagtattcgaagttattcgaacagttctaatccagacgttaaga
1 M T R L R S I K T S G W K E Y S K L F E T V L I Q T L R
4382 ctcacgcatttgggatga 4399
29 L T H L G *
dp1ORF260
24746 gtgacctacttctcctcaatcgccggtactggaggcaagcaagctcaagtcacttccatttcaggaaacttcaacttcttccag
1 V T L L P Q S A V L E A S K L K S L P F Q E T S T S F Q
24830 cggctgaatattatttag 24847
29 R L N I I *
dp1ORF261
288 atgaattcacttccctttgccctaaaacaggacagcctgacttcgcgaatgttttcattagttacattccaaacgaaaagatgg
1 M N S L P F A L K Q D S L T S R M F S L V T F Q T K R W
372 ttgaatcctaaatcattga 389
29 L N L N H *
dp1ORF262
9408 atgcctattcaactccaggcggaaaagatgtggaagcatgctgtgagtcagtttaaattagaaaaggtgactaccttgacg
1 M P I Q L Q A E R C G S M L V Q F D L N L E K V T T L T
9492 aaaacgggtgcatcattga 9509
29 K T V H H *
dp1ORF263
27052 atgaaaatttttagcatcgagttctttcgaagttttcgaataatttcccttcacctgtttgatagttggttcatctagacctttt
1 M K I L A S S S F E V F E I I S F T C L I V G S S R P F
26968 aacaagtcttctaattga 26951
29 N K S S N *
dp1ORF264
6139 gtgaatagttacaaggcgggtctaatacgcctcaggattttctgctgtaggatagccgcacatcttcaaaactcaattgagtcgaagc
1 V N S T R R S N T L R I S A V G I A A S S S N S I E S S
6055 tgtgaaacgtcttcataa 6038
29 C E T S S *
dp1ORF265
4801 gtgaataaagtcaagcgtttttgtataaaaagttcatttttttttaaaaaataagagcgaaaagctcttatctaaaatagtc
1 V N K V K R F C I K S S F F P K K N K S E K L L S K I V
4717 gacgttgacgatttttaa 4700
29 D V D D F *
dp1ORF266
50220 atgcccgttcttccaagcagttgcaagcattttatcaatagtcacgacttaccttgtccaggctcgagccattatgacaatcaa
1 M P V L P S S C K H F I N S P R L T L S R S S H Y D N Q
50136 atcctcaccaggaagtaa 50119
29 I L T R K *
dp1ORF267
47367 atggtaacaggtctgttctaggttcaggaagaacaaacgggaagtgaatgttattttcttcagcgaagttcttttgcttcatacca
1 M V K V C S R F R K N K R E V N V I F F S E V F C F I P
47283 aacattaatcgtagatag 47266
29 N I N R R *
dp1ORF268

397

12621 atgtcaatttcgggtcttgtgcttgacaatggattcaactactgatgcgtcaacctttttcaatcgcgacagcttgcattca
1 M S I S V L C L T M D S T T D A S T F F N R D S L S N S
12537 ttgtcaattctagagtaa 12520
29 L S I L E *
dp1ORF269
53834 gtgaatagtatcgagtcctcagtttctacgtcaatagaacctattccgtcttcaatcattttgtctacatactgctcgagttt
1 V N S I E S I S F Y V N R T Y S V F N H F V Y I L L E F
53750 tgccttcctcagtgattaa 53733
29 C F L S D *
dp1ORF270
50792 atgatttttcgggtcttcgccatatcggtttttaacgacagatagttcaagtatgccggatttttcgtcacgcttcatacgagata
1 M I F R S S P Y R F L T T D S S S M P D F S S R F I A I
50708 actctgctagcattttga 50691
29 T L L A F *
dp1ORF271
19739 atgaggctgctttgctttatcttcgttacgtattgaccgacttcctactcggaaccttcctacaagaattcatacctcaaag
1 M R L L C F I F V T V L T D F L L A N L P T R I H T S K
19655 gctttttgtcagccttag 19638
29 A F C Q P *
dp1ORF272
1556 gtgggtcaagtctgtcaatgaatgtacctgcgattttcttgacgtgataaaagtcaacaacctcccttgactcgaaccgtggtc
1 V V K S V N E C T C D F L D V I K V N N H P L T R T V V
1472 ataagttccgctgctaa 1455
29 I S S A C *
dp1ORF273
56256 atggatttccattaggactgagtcctcttgggaattggaacggttgcatatatagatattccgtcagccgtactaggccaagttct
1 M D F I R T E S S W N W N G C I Y R Y S V S R T R P S S
56340 agttcagttttatcttgacgtcaattgcttcgagatatttgaaaaagtagtcaggaaaattcctgattatcttgacgtcaattgc
29 S S V Y L A V N C F E I F E K V V R K I P D Y L A V N C
56424 ttcgagatatttgaaaaagtagtcaggaaaattcctgattatttttttcaaaaaacgcttga 56486
57 F E I F E K V V R K I P D Y F F Y K N A *

398

Table 31

Query= sid|114822|lan|dp1ORF001 Phage dp1 ORF|36698-40390|2
(1230 letters)

>gi|928828 (L44593) ORF1904; putative [Lactococcus lactis phage BK5-T]
Length = 1904

Score = 427 bits (1086), Expect = e-118
Identities = 226/475 (47%), Positives = 281/475 (58%), Gaps = 45/475 (9%)

Query: 395 AESGKYIGVLNTNKKPSELVPDDFTWIRLEGPKGDAGLPGAPGRDGVDPGKSGVGIAD 454
A+ YIG + P D+TW + +G+ G GA G+DGV GK GVGI
Sbjct: 820 ADYPSYIGQYTDIFIQYDSAKPSDYTWSLI---RGNDGKDGATGKDG---AGKDGVGIKT 873

Query: 455 TAITYAVSVSGTQEPENGWSEQVPELIKGRFLWTKTFWRYTDGSSETGYSVAYIGQDGN 514
T ITYA+S SGT +P GW+ QVP L+KG++LWTKT W YTD S ETGYSV YI +DGN+
Sbjct: 874 TVITYALSSSGTDKPNWTGWSQVPTLVKGQYLWTKTVWYTTDSSSETGYSVTYIAKDGN 933

Query: 515 GKDGIAKDGVGIAATEVMYASSPSATEAPAGWSTQVPTVPGGQYLWTRTRWRYTDQTD 574
G DGIAGKDGVGI T + YA S T APA GW++QVP VP GQ+LWT+T W YTD T
Sbjct: 934 GNDGIAKDGVGIKKTTITYAVGTSGTTAPASGWSQVPNVPAGQLWTKTVWYTTDNTS 993

Query: 575 EIGYSVSRMGEQGPKG DAGR---DGIAGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVP 630
E GYSV+ MG +G KGD G +GIAGK+G G+K+T+++Y SP + P G W++ VP
Sbjct: 994 ETGYSVAMMGVKGDKGDPGNNGTNGIAGKDGKIKATAITYQASPNGTAPTGTWSASVP 1053

Query: 631 SLIKGQYLWTRTIWYTDSTTETGYQKTYIPKDGNDGKNGIAGKDGVGIKSTTITYAGST 690
+ KG +LWTRTIWYTD+TTETGY Y+ +GN+G +G GKDG GIK+TTITYAGST
Sbjct: 1054 PVAKGSFLWTRTIWYTDNTTETGYAVAYMGNGNGHGDGFPKGDTGIKTTITYAGST 1113

Query: 691 SGTVAPTSNTSAIPNVQPGFFLWTKTVWNYTDDTSETGYSVSKIGETXXXXXXXXXXXX 750
SGT P + WTS +P V G +LWTKTV W YTD+TSETGYSV+ +G
Sbjct: 1114 SGTTPPNNGWTSTVPTVAEGNYLWTKTVWYTTDNTSETGYSVAMMG-----VKGDKGD 1167

Query: 751 XXXXXXXXXXXXADGRS-QYTHLAFSNSPNGEFSGHTDSGRAYVGGYQDFNPVHSDPAAYT 809
DG+ + T + + SPNG A G + P +K +T
Sbjct: 1168 GNGTNGIAGKDGKIKATAITYQASPNGT-----TAPTGTWSASVPPVAKGSFLW 1219

Query: 810 WTKN-----KGNDGAQGIPIGKPGADGKTNYPHIAYASSADGS 846
T W GN+G G PGK G KT I YA S G+
Sbjct: 1220 RTIWTYTDNTTETGYAVAYMGNGNGHGDGFPKGDTGIKTT--TITYAGSTSGT 1272

Score = 396 bits (1007), Expect = e-109
Identities = 208/449 (46%), Positives = 260/449 (57%), Gaps = 42/449 (9%)

Query: 421 IRLEGPKGDAGLPGAPGRDGVDPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480
+ + G KGD G PG +G +G+ CK G GI TAITY S +GT P WS VP +
Sbjct: 1155 VAMMGVKGDKG---DPGNNGTNGIAGKDGKIKATAITYQASPNGTAPTGTWSASVPPV 1211

Query: 481 IKGRFLWTKTFWRYTDGSSETGYSVAYIGQDGN SGKDGIAKDGVGIAATEVMYASSPSA 540
KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S
Sbjct: 1212 AKGSFLWTRTIWYTTDNTTETGYAVAYMGNGNGHGDGFPKGDTGIKTTITYAGSTSG 1271

Query: 541 TEAPAGWSTQVPTVPGGQYLWTRTRWRYTDQTD EIGYSVSRMGEQGPKG DAGR---DGI 597
T P GW++ VPTV G YLWT+T W YTD T E GYSV+ MG +G KGD G +GI
Sbjct: 1272 TTPPNNGWTSTVPTVAEGNYLWTKTVWYTTDNTSETGYSVAMMGVKGDKGDPGNNGTNGI 1331

Query: 598 AGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVPSLIKQYLWTRTIWYTDSTTETGYQ 656
AGK+G G+K+T+++Y SP + P G W++ VP + KG +LWTRTIWYTD+TTETGY
Sbjct: 1332 AGKDGKIKATAITYQASPNGTAPTGTWSASVPPVAKGSFLWTRTIWYTTDNTTETGYA 1391

Query: 657 KTYIPKDGNDGKNGIAGKDGVGIKSTTITYAGSTSGTVAPTSNTSAIPNVQPGFFLWTK 716
Y+ +GN+G +G GKDG GIK+TTITYAGSTSGT P + WTS +P V G +LWTK
Sbjct: 1392 VAYMGNGNGHGDGFPKGDTGIKTTITYAGSTSGTTPPNNGWTSTVPTVAEGNYLWTK 1451

Query: 717 TVWNYTDDTSETGYSVSKIGETXXXXXXXXXXXXXXXXXXXXADGRS-QYTHLAFSNS 775
TVW YTD+TSETGYSV+ +G DG+ + T + + S
Sbjct: 1452 TVWYTTDNTSETGYSVAMMG-----VKGDKGDGPNNGTNGIAGKDGKIKATAITYQAS 1505

399

Query: 776 PNREGFSHTDSGRAYVGGYQDFNPFVHSDFAAYTWIKW-----KGND 817
PNG A G + P +K +T T W GN+
Sbjct: 1506 PNGT-----TAPTGTWSASVPPVAKGSFLWTRTIWYTDNTTETGYAVAYMGNGNN 1557

Query: 818 GAQGI PGKPGADGKTNYPHIAASSADGS 846
G G PGK G KT I YA S G+
Sbjct: 1558 GHDGFPKDGITGIKT--TITYAGSTSGT 1584

Score = 384 bits (977), Expect = e-105
Identities = 179/322 (55%), Positives = 222/322 (68%), Gaps = 7/322 (2%)

Query: 421 IRLEGPKGADAGLPGAPGRDGVDPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480
+ + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP +
Sbjct: 1311 VAMMGVKGDKG---DPGNNGTNGIAGKDGKGIKATAITYQASPNGTAPTGTWSASVPPV 1367

Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGN SGKDG IAGKDGVGIAATEVMYASSPSA 540
KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S
Sbjct: 1368 AKGSFLWTRTIWYTDNTTETGYAVAYMGNGNHDGFPKDGITGIKTITITYAGSTSG 1427

Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQDEIGYSVSRMGEQGPKGADAGR---DGI 597
T P GW++ VPTV G YLWT+T W YTD T E GYSV+ MG +G KGD G +GI
Sbjct: 1428 TTPPNNGWTSTVPTVAEGNYLWTKTVWYTDNTSETGYSVAMMGVKGDKGDPGNNGTNGI 1487

Query: 598 AGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVPSLIKQYLWTRTIWYTDSTTETGYQ 656
AGK+G G+K+T++Y SP + P G W++ VP + KG +LWTRTIWYTD+TTETGY
Sbjct: 1488 AGKDGKGIKATAITYQASPNGTAPTGTWSASVPPVAKGSFLWTRTIWYTDNTTETGYA 1547

Query: 657 KTYIPKDGNDGKNGIAGKDGVG IKTITITYAGSTSGTVAPTSNWTSAIPNVQPGFFLWTK 716
Y+ +GN+G +G GKDG GIK+TTITYAGSTSGT P + WTS +P V G +LWTK
Sbjct: 1548 VAYMGINGNNGHDGFPKDGITGIKTITITYAGSTSGTTPPNNGWTSTVPTVAEGNYLWTK 1607

Query: 717 TVWNYTDDTSETGYSVSKIGET 738
TVW YTD++ ETGYSV K+G T
Sbjct: 1608 TVWAYTDNSFETGYSVGKMGNT 1629

Score = 201 bits (507), Expect = 2e-50
Identities = 121/297 (40%), Positives = 156/297 (51%), Gaps = 19/297 (6%)

Query: 421 IRLEGPKGADAGLPGAPGRDGVDPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480
+ + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP +
Sbjct: 1467 VAMMGVKGDKG---DPGNNGTNGIAGKDGKGIKATAITYQASPNGTAPTGTWSASVPPV 1523

Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGN SGKDG IAGKDGVGIAATEVMYASSPSA 540
KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S
Sbjct: 1524 AKGSFLWTRTIWYTDNTTETGYAVAYMGNGNHDGFPKDGITGIKTITITYAGSTSG 1583

Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQDEIGYSVSRMGEQGPKGADAGRDIAGK 600
T P GW++ VPTV G YLWT+T W YTD + E GYSV +MG GP AG +G GK
Sbjct: 1584 TTPPNNGWTSTVPTVAEGNYLWTKTVWAYTDNSFETGYSVGKMGNTGP---AGSNGNPGK 1640

Query: 601 NGIGLKSTSVSYGISPTDSAIPGVWASQVPSLIKQ-YLWTRTIWYTDSTTE--TGYQK 657
+ T+ G++ S + + ++ G +Y W W + G
Sbjct: 1641 VVSDTEPTTKFKGLTWKYSVGVDMPLGNGTKILAGTEYYWNGNWNALYEINAHNINGDNL 1700

Query: 658 TYIPKDGNDGK-NGIAGKDGVG IKTITITYAGS-----TSGTVAPTSNWTSAIPNVQ 708
+ DGK I G +GV + T T GS +S + T N T A I N Q
Sbjct: 1701 SVTNGTFKDGKIESIWGSNGV---NGTTIEGSHLQIHSSDSTNTTEN-TLAIDNRQ 1753

Query= sid|114823|lan|dp1ORF002 Phage dp1 ORF|32386-35835|1
(1149 letters)

>dbj|BAA31888| (AB009866) orf 15 [bacteriophage phi PVL]
Length = 694

Score = 280 bits (709), Expect = 3e-74
Identities = 157/465 (33%), Positives = 257/465 (54%), Gaps = 28/465 (6%)

Query: 40 QIGSALTGLGKGLTAVTLPLMGFAAASIKVGNFQAQMSRVQAIAGATAEELGRMKTQA 99
+IG+++ +G+ +T VT P++ A + K G EF M +V+A +GAT EE +K +A
Sbjct: 151 EIGNSMKNVGRNMTMYVTAPVAVGFAVAAKKGIEFDDSMRKVKATSGATGEEFRAALKKA 210

400

Query: 100 IDLGAKTAFSAKEAAQGMENLASAGFQVNEIMDAMPVGLDLXXXXXXXXXXXXMASSL 159
++GA T FSA +++A+ + +A AG+ ++M+ + GV+DL + L
Sbjct: 211 REMGATTKFSASDSABALNYMALAGWDSKQMMEGLSGVMDLAAASGEELGAVSDIVTDGL 270

Query: 160 RAFGLEAQAGHVADVFARAAADTNAETSDMAEAMKYVAPVAHSMGLSLEETAASIGIMA 219
AFGL+A +GH+ADV A+ ++ N + + EA KYVAPVA ++G ++E+T+ +IG+M+
Sbjct: 271 TAFGLKAKDSGHLADVLQATSSKANTDVRGLGEAFKYVAPVAGALGYTIEDTSIAIGLMS 330

Query: 220 DAGIKGSQAGTTLRGALSRIAKPTKAMVKSMQELGVSYFDANGNMIPLREQIAQLKTATA 279
+AGIKG +AGT LR + ++ PT+AM M+ LG+S D+NG MIP+R+ + QL+
Sbjct: 331 NAGIKGEKAGTALRTMFTNLSSPTRAMGNEMERLIGISITDSNGKMI PMRKLDDQLREKFK 390

Query: 280 GLTQEERNRHLVTLYGQNSLSGMLALLDAGPEKLDKMTNALVNSDGAAKEMAETMQDNLA 339
L+++++ T++G+ ++SG LA+++A E K+T ++ +S GA+K MA+TM+ L
Sbjct: 391 HLSKDQQAASSAATIFGKEAMSGALAIINASDEDEYQKLTKSIDSSTGASKRMADTMESGLG 450

Query: 340 SKIEQMGGAFESVAIIVQQILEPALAKIVCAITKVLEAFVNMSPIGQKMMVIFAGMVAAL 399
K+ + E +A+ + +EPAL IV A +KV+ + Q VV F VA L
Sbjct: 451 GKLRTRLSQLEELALTIYDRIEPALKIIVSAPSKVVTWTKLPTSQILAVVGFGLFVAVL 510

Query: 400 GPLLLIAGM-----VMTTIVKLRIAIQFLGPAFMGTMGTIAGVIAIF----- 441
GPL+ + G+ MT + L I + F IA ++ +F
Sbjct: 511 GPLVFMFGLFISVMGNAMTVLGPLLINVNKASGLFAFLRTKIASLVKLFPIILGVSISSLT 570

Query: 442 -----YALVAV---FMIAYTKSERFRNFINS LAPAIKAGFGGA 476
ALV + F AY +SE FRN +N + F A
Sbjct: 571 LPITLIVGALVGIGIAFYQAYKRSETFRNIVNQAISGVANAFKAA 615

Query= sid|114824|lan|dp1ORF003 Phage dp1 ORF|53538-55877|3
(779 letters)

>sp|P43741|DPOL_HAEIN DNA POLYMERASE I (POL I) >gi|1074025|pir||E64098 DNA polymerase I
(polA) homolog - Haemophilus influenzae (strain Rd KW20)
>gi|1573871 (U32767) DNA polymerase I (polA)
[Haemophilus influenzae Rd]
Length = 930

Score = 191 bits (481), Expect = 1e-47
Identities = 148/553 (26%), Positives = 262/553 (46%), Gaps = 60/553 (10%)

Query: 63 RLELITEAKLEQYVDKMIEDGIGSIDVETDGLDTIHDELAVCLYSPQKGIYAPVNVH 122
+ E + +A L +++K+ + ++D ETD LD + L G+ + + Y P+
Sbjct: 333 KYETLLTQADLTRNIEKLNAAKLIADVTETDSDLVMSANLVGISFALENGEAAAYLPQLD 392

Query: 123 SNMTKMKRIKNQISPEFMKKMLQRIVDSGIPVIYHNSKFDMSKIYWRIGVKMNEPAWDTYL 182
++ + +K +L+ + I I N KFD +SI+ R G+++ +DT L
Sbjct: 393 YLDAPKTEKSTALAAIKPILE---NPNHKGIGQNIKFD-ESIPARHGIELQGVFDTML 448

Query: 183 AAMLLNENESHSLKSLHSKYVRNEENAFAKFNDFKGPFPFLIPPDVAYMYAAYDPLQT 242
+ LN H++ L +Y+ +E A + + F+ IP + A YAA D T
Sbjct: 449 LSYTLNSTGRHNMDDAKRYLGHETIAFESLAGKGSQLTFFNQIPLEQATEYAADADVT 508

Query: 243 FELYEQEQYLTPGTEQCEEYNLEKVSWSVLHNIEMPLIKVLFDMVEYGVLDQDKLAEIR 302
+L + L E Y +E+PL+ VL ME GV +D D L
Sbjct: 509 MKLQQALWLKLQEPTLVELYK-----TMELPLLHVLSRMERTGVLIDSALFMQS 559

Query: 303 EQFTANMNEAEQEQQLVSEWQPEIEELRQTNFQSYQKLEMDARGRTVSISSTQLAIL 362
+ + + E++ L + +++S QL +
Sbjct: 560 NEIASRLTALEKQAYALAGQ-----PFNLASTKQLQEI 592

Query: 363 FYDIMGLKSPERDKPRG---TGESIVEH--FDNDISXXXXXXXXXXXXVSTYTT-LDQHL 416
+D + L ++ P+G T E ++E + +++ STYT L Q +
Sbjct: 593 LFDKLELFPVLQKT-PKGAPSTNEEVLEELSYSHELPKILVKHRLSKLSTYTDKLPQMV 651

Query: 417 AKPDNRHITTFKQYGAKTGRMSSSENPNLQNIPIRGE-GAVVRQIFAASEGHYIIGSDYSQ 475
R+HT++ Q TGR+SS +PNLQNIPI R E G +RQ F A EG+ I+ +DYSQ
Sbjct: 652 NSQTGRVHTSYHQAATATGRLSSSDPNLQNIPIRNEEGRHRIQAFIAREGYSIVAADYSQ 711

Query: 476 QEPRSLAELSGDESMRHAYEQNLDSLVSIGSKLYGVFYECELEFPDGTNKEGKLRRNS 535
E R +A LSGD+ + +A+ Q D++ +++++GV +E T+++ R +
Sbjct: 712 IELRIMAHLSGDQGLINAFSQGKDIHRSTAAEIFGVSLDE-----VTSEQ-----RRN 759

Query: 536 VKSVLLGLMYGRGANSIAEQMNVSVKEANKVIEDFFTEFPKVADYIIFVQQQAQDLGYVQ 595
K++ GL+YG A ++ Q+ +S +A K ++ +F +P V ++ +++++A+ GYV+

401

Sbjct: 760 AKAINFGLIYGMSAFGLSRQLGISRADAQKYMDLYFQRYPSVQQFMTDIREKAKAQGYVE 819

Query: 596 TATGRRRRRLPDMS 608

T GRR LPD++

Sbjct: 820 TLFGRRLYLPDIN 832

Score = 46.9 bits (109), Expect = 5e-04

Identities = 34/123 (27%), Positives = 66/123 (53%), Gaps = 16/123 (13%)

Query: 663 EIKDQAKAEGI-----LIKDNNGKIADAQRQCLNSVIQGTAAADMTKYAMIKV 709

+I+++AKA+G + N + A+R +N+ +QGTAA+ K AMIK+

Sbjct: 807 DIREKAKAQGYVETLFGRRLLYLPDINSSNAMRRKGAERVAINAPMQGTAADIKRAMIKL 866

Query: 710 HNDDELKELGFLMIPVHDELLGEVPIKNAKGAERLTEVMIEAAKDIISLPMKCDPSIV 769

++ + +++ VHDEL+ EV + E++ + M EAA +++ +P+ + +

Sbjct: 867 -DEVIRHDPDIEMIMQVHDELVFEVRSEKVAFFREQIKQHM-EAAAELV-VPLIVEVGVG 923

Query: 770 ERW 772

+ W

Sbjct: 924 QNW 926

Query= sid|114825|lan|dp1ORF004 Phage dp1 ORF|40401-42440|3
(679 letters)

>emb|CAB07981| (Z93946) hypothetical protein [bacteriophage Dp-1]
Length = 532

Score = 1011 bits (2585), Expect = 0.0

Identities = 497/499 (99%), Positives = 498/499 (99%)

Query: 1 MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLN 60

MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLN

Sbjct: 1 MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLN 60

Query: 61 SSVHSSHPDYDTSGEVTLASGEVTVPHNSDGTKTMSVWASFDPMNGVHGNTISTNYTL 120

SSVHSSHPDYDTSGEVTLASGEVTVPHNSDGTKTMSVWASFDPMNGVHGNTISTNYTL

Sbjct: 61 SSVHSSHPDYDTSGEVTLASGEVTVPHNSDGTKTMSVWASFDPMNGVHGNTISTNYTL 120

Query: 121 DSIPRSTQISSFEGNRLGSLHTVIFNRKVNSTHQVWYRVFGSDWIDLGNHTTSVSFT 180

DSIPRSTQISSFEGNRLGSLHTVIFNRKVNSTHQVWYRVFGSDWIDLGNHTTSVSFT

Sbjct: 121 DSIPRSTQISSFEGNRLGSLHTVIFNRKVNSTHQVWYRVFGSDWIDLGNHTTSVSFT 180

Query: 181 PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYNSGWRFNIPDSVRPTFSGISLVDTT 240

PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYNSGWRFNIPDSVRPTFSGISLVDTT

Sbjct: 181 PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYNSGWRFNIPDSVRPTFSGISLVDTT 240

Query: 241 SAVRQILTGNNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGKGLGMMNF 300

SAVRQILTGNNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGKGLGMMNF

Sbjct: 241 SAVRQILTGNNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGKGLGMMNF 300

Query: 301 NGSATVRAWVTDTRGKQSNVQDVSINVIEYGPSINFSVQRTQNPAAIQALRNKAVAPI 360

NGSATVRAWVTDTRGKQSNVQDVSINVIEYGPSINFSVQRTQNPAAIQALRNKAVAPI

Sbjct: 301 NGSATVRAWVTDTRGKQSNVQDVSINVIEYGPSINFSVQRTQNPAAIQALRNKAVAPI 360

Query: 361 TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGFTTISLMTNSSANLAGNYGPKSYIV 420

TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGFTTISL+TNSSANLAGNYGPKSYIV

Sbjct: 361 TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGFTTISLMTNSSANLAGNYGPKSYIV 420

Query: 421 KAKIQDRFTSTEPSATVATESVVLNYDKDGRIGVGKVVEQKAGSIDAAGDIYAGGRQVQ 480

KAKIQDRFTSTEPSATV TESVVLNYDKDGRIGVGKVVEQKAGSIDAAGDIYAGGRQVQ

Sbjct: 421 KAKIQDRFTSTEPSATVPTESVVLNYDKDGRIGVGKVVEQKAGSIDAAGDIYAGGRQVQ 480

Query: 481 QFQLTDNNGALNRGQYNDV 499

QFQLTDNNGALNRGQYNDV

Sbjct: 481 QFQLTDNNGALNRGQYNDV 499

Query= sid|114827|lan|dp1ORF006 Phage dp1 ORF|45296-46987|2
(563 letters)

>gb|AAD18987| (AE001666) SWI/SNF family helicase_2 [Chlamydia pneumoniae]
Length = 1166

Score = 171 bits (429), Expect = 1e-41

Identities = 150/522 (28%), Positives = 254/522 (47%), Gaps = 55/522 (10%)

402

Query: 46 SSNNPE-LPYKYFNNVIDALDEWELHIFGELDKDVQDYIDSRNRIASSSNEQSFKTTTPF 104
S + FE LP + ++ + L E + I GE++ D QD + T

Sbjct: 659 SLDQFEALPVNF--SMSERLIEIQKQIRGEIEFDQD-----VPQIQATLRSYQTEG 709

Query: 105 AHQVECFEYAEHPCFLLGDEQGLGKTQQAIDIAVSRAKSPKH--CLIVCCISGLKWNWA 162
H +E + H +L D+ GLGKT QAI IAV++ K C ++ C + L +NW

Sbjct: 710 VHWLE--RLRKMHNLGILADDMLGKTLQAI-IAVTQSKLEKSGCSLIVCPTSLVYNWK 766

Query: 163 KEVGIHSNESAHILGSRVTKDGLVIDGV-SKRAEDLLGGHDEFFLITNIETLRDAVFIK 221
+E + E LVIDGV S+R + L D IT+ L+ V

Sbjct: 767 EEFKRFNPEFR-----TLVIDGVPSQRRKQLTALADRDVAITSYNLLQKDV--- 812

Query: 222 YLNELTKSGEIGMVIIDEIHKCKNPSSKQASIQKLSYKMGTLGTPLMNNPIDVFNVM 281
EL KS V++DE H KN +++ S++ +QS +++ LTGTP+ N+ +++++

Sbjct: 813 ---ELYKSFRFDYVVLDEAHHIKNRTRNAKSVKMIQSDHRLILTGTPIENSLEELWSLF 869

Query: 282 KWLGAEHHTLTQPKERYCIVDQFNQITGYR-----NLAEELVNDYMLRRTKEEVL-DL 335
+L L +R+ V ++ + Y N+ L++ V+ ++LRR KE+VL DL

Sbjct: 870 DFLMPG---LLSSYDRF--VGKYIRTGNMGKNADNMVALKKKVSFFILRRMKEDVLKDL 924

Query: 336 PEKIRVTEYVDMNSKQSKIY-----KEVLTKLVQEIDKVKLMPNPLAETIRLRQATGN 388
P + + + Q ++Y K+ L++LV++ ++ + LA RL+Q +

Sbjct: 925 PVPSEILYHCHLTESQKELYQSYAASAKQELSRVLKQEGFERIHHVLA TLRLKQICCH 984

Query: 389 PSILTTQDVK---SCKFERCIEIVEECIQGKSCVIFSNEKVVIEPLAKIL-SKTVKCNL 444
P+I + S K++ +++ + G V+FS + K++ + K L S+ +

Sbjct: 985 PAIFAKDAPEPGDSAKYDMLMLLLSSLVDSGHKTVVFSQYTKMLGIKKDLESRGIPFVY 1044

Query: 445 VTGETADKFNEIEEFMNRKASVILGTIGALGTGTLTKADTVIFLDSWPWTRAEKDQAE 504
+ G T ++ + + +F V L ++ A GTG L ADTVI D W A ++QA D

Sbjct: 1045 LDGSTKNRLDLVNQFNEDPSLLVFLISLKAGGTGLNLVGGADTVIHYDMWNPVAVENQATD 1104

Query: 505 RCHRIGAKSSVTIYTLVAKGTVDERIEDLIERKGELADYIVD 546
R HRIG SV+ Y LV T++E+I L RK L +++

Sbjct: 1105 RVHRIGQSRVSSYKLVTLNTIEEKILTQNRKSLVKKVIN 1146

Query= sid|114828|lan|dp1ORF007 Phage dp1 ORF|22230-23621|3
(463 letters)

>gi|2444105 (U88974) ORF26 (Streptococcus thermophilus temperate bacteriophage
O1205)
Length = 411

Score = 88.9 bits (217), Expect = 7e-17
Identities = 80/315 (25%), Positives = 133/315 (41%), Gaps = 48/315 (15%)

Query: 139 QGVTLAGIFCDEVALMPESFVNQATGRCSVTGSKMWFSCNPANPNHYFKKNWIDKQVEKR 198
+G T G + +E +L E + RCS G+++ + NP NPNH+ +++I K + +

Sbjct: 121 RGFTAFGAYVNEASLANELVFKEIISRCSGDARVVWDSNPDNPNHWLNRDYIGKN-DGK 179

Query: 199 ILYLHFTMDDNPSTL---DSIKRREKMYAGVFRKRFLGLWVTADGLVYSMFNEEQHV 254
I+ F +DDN L+ DSIK K G F R ILGLW A+G +Y+ ++ + HV

Sbjct: 180 IIDFSFKLDDNTFLSKRYIDSIAKATPK---GKPYDRDILGLWTVAGAIYADYDSKIHV 236

Query: 255 KKLNIEFDRLFVAGDFGIYNATTFGLYGFSKRHKRYHLIESYHSGREAEQLTEADVNS 314
E R P D+G + + + G ++L++ +E + + +A

Sbjct: 237 VDELPEMKRYFGGIDWGYTHYSIVIVG-EGVDNNFYLVDGVAAQFKEIDWVVEQA---- 291

Query: 315 NIQFSSVLQKTTKEYANDLVDMIRGKQIEYIILDPSASAMIVELQXHPYIAR---KNIP 371
+K T Y N + + ++AR + I

Sbjct: 292 -----RKLGTGIYGN-----IPFYADSRPEHVARFENEGFDI 323

Query: 372 IPARNVDTLGISFAELLAENRFTLDPSNT-HDIDEYYAYSWDSKASQTGEDRVIKEHDH 430
+ A V GI A+L E + + DE Y Y W ++ +D +KE D

Sbjct: 324 MNANKSVIAGIELIAKLFKEKKLYVKRGFVPRFFDEIYQYRWKENST---KDEPLKEFDD 380

Query: 431 CMDRNRVYACLTDALI 445
+D RYA +D +I

Sbjct: 381 VLDSVRYAIYSDYVI 395

Query= sid|114829|lan|dp1ORF008 Phage dp1 ORF|49624-50961|1
(445 letters)

>gb|AAD19901| (AF100420) DnaB replication fork helicase (Thermus aquaticus)

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Length = 444

Score = 67.5 bits (162), Expect = 2e-10
 Identities = 69/248 (27%), Positives = 111/248 (43%), Gaps = 14/248 (5%)

Query: 147 GERLGISTGFEXXXXXXXXXXXXXXXXXXIVIMARPGQGS-WTIDKMLATAWKNHVDVLLYS 205
 GE G+ TGF+ I I ARP GK+ + + A K G V +YS
 Sbjct: 178 GEVAGVRTGFKELDLIGTLGPGSLNI-IAARPAMGKTAFALTIAQNAALKEGVGVGIYS 236

Query: 206 GEMSEMVGARIDTILSNVSINSITKGIWNHDFEYEDHIQAMTEANSLVVTPFMIG 265
 EM Q+ R+ + + N+ G D F+ D ++EA + TP +
 Sbjct: 237 LEMPAQLTLRMMCSEARIDMNRVRLGQLTDRDFSRSLVDVASRLSEAP-IYIDTPDLTL 295

Query: 266 GKNLTPAILDSMISKYRPSVVGIDQLSLMS--ESYPSREQKRIQYANITMDLYKISAKYG 323
 + A ++S+ + ++ ID L LMS S S E ++ + A I+ L ++ + G
 Sbjct: 296 ME--VRARRLVSQNQVGLIIDIYQLMSGPGSGKSGENRQOEIAAISRGLKALARELG 353

Query: 324 IPIVLNVQAGRSKTEGAESMELEHIAESDGVQNASRVIAMKRD-----EKSGLILEL 376
 IPI+ Q R+ + + L + ES + Q+A V+ + RD EK+GI E+
 Sbjct: 354 IPIIALSQLSRAVEARPKNRPMLSDLRESGSIEQDADLVMFYRDEYYNPHSEKAGIAEI 413

Query: 377 SVVKRYG 384
 V K R G
 Sbjct: 414 IVGKQRNG 421

Query= sid|114831|lan|dp1ORF010 Phage dp1 ORF|8699-9859|2
 (386 letters)

>gi|2760912 (AF037258) RecA protein [Chlorobium tepidum]
 Length = 346

Score = 133 bits (331), Expect = 2e-30
 Identities = 99/340 (29%), Positives = 164/340 (48%), Gaps = 66/340 (19%)

Query: 44 GGLPRKRVEFFGPSSSGKTTSDIVKNAQMVFXXXXXXXXXXXXXXXXXXARASKASKT 103
 GGLPR RV E +GPSSSGKTT AL + AQ
 Sbjct: 67 GGLPRGRVTEIYGPSSSGKTTLALHAIAEAQ-----KNG 100

Query: 104 AVKELEMQLDSLQEPKIVYLDLENTLDEWAKKIGVDVDNINIVRPEMNSAEELQYVL 163
 + L +D E+ D +A+K+GVD++ + + +PE S E+ L V
 Sbjct: 101 GIAAL-----VDAEHAFDPTYARKLGVDINALLVSQPE--SGEQALSIVE 143

Query: 164 DIFETGEVGLVLDLSPYMSQNLIDEELTKKAYAGISAPLTFESRKVTPLLTRYNAIFL 223
 + +G V ++V+DS+ +V Q ++ E+ + +++ RK+T +++ +++ L
 Sbjct: 144 TLVRSQAVDIIIVDSVAALVPQAELEGEMGDSVVGLQARLMSQALRKLTAISKSSSVCL 203

Query: 224 GINQIREDMNSQYNA-YSTPGGKMKHACAVRLKFRKGDYLDENGASLRTARNPAGNVV 282
 INQ+R+ + Y + +T GKG K +VRL RK + ++G L GN
 Sbjct: 204 FINQLRDKIGVMYGSPETTTGKALKFYSSVRLDIRKIAQI-KDGEELV-----GNRT 255

Query: 283 ESFVEKTKAFKPDRLVSYTSLYHDIQIENDLVDVAVEFGVIQKAGAWFSIVDLETGEI 342
 + V K K P K + + Y +GI + +L+D+AVEFG+I+K+GAWFS + G
 Sbjct: 256 KVKVVKQKV-APPFKTAEFDILYGEGISVLGELIDLAVEFGIIKSGAWFSYGTEKLG-- 312

Query: 343 MTDEDEEPLKFQKANKLVRRFKEDDYLFDMVMTAVHEIIT 382
 QG+ N+ + KED+ L + + V +++T
 Sbjct: 313 -----QRENVKLLKEDETLRNTIRQQVRDMLT 341

Query= sid|114832|lan|dp1ORF011 Phage dp1 ORF|28017-29096|3
 (359 letters)

>gi|2444110 (U88974) ORF31 [Streptococcus thermophilus temperate bacteriophage
 O1205]
 Length = 348

Score = 187 bits (469), Expect = 1e-46
 Identities = 118/358 (32%), Positives = 187/358 (51%), Gaps = 21/358 (5%)

Query: 3 IYDYNAGEIASYIQALPSNALQYLGPITLFPNAQQTGTDISWLKGANLPTVIQPSNYDA 62
 IYD + A IA Y AL N LG ++FP +Q GT +S++KGA+ V ++ + +D
 Sbjct: 4 IYDKVTASNIAGYFNALQENVSTLGSIFPARKQLGTLKLSYIKGASGQSVALKAAAFDT 63

Query: 63 KASLRERAGFSKQATEMAFFRESMRLGEKDRQNLQMLLNQSSA-LAQPLITQLYNDTKNL 121

404

++R+R +M FF+E+M + E DRQ L ++ + +A L ++ ++ND L
 Sbjct: 64 NVTIRDRVSAEMHDEQMPFFKEAMLVKENDRQQLNLVKDSGNAVLVNTIVAGIFNDNLTL 123

Query: 122 VDGVEAQAEYMRMQLLYGKFTVKSTNSEAQYTYDYNMDAKQYAVTKKWTNPAESDPIA 181
 V+G A+ E MRMQ+L GK S Y D K+Q V+K W P + P+A
 Sbjct: 124 VNGAPARLEAMRMQVLATGKIAFTSDGVNKDIDYGVKPDHKKQ--VSKSWAEPG-ATPLA 180

Query: 182 DILAAMDDIENRTGVRPTRMVLNRNTYNQMTKSDSIKKAL-AIGVQGSWENFLLASDAE 240
 D+ A+ + G+ P R V+N T+ + K+ S K + + GS + ++ E
 Sbjct: 181 DLEDAI-ETARELGLNPERAVMNAKTFGLIRKAASTVKVIKPLAGDGS----AVTKAELE 235

Query: 241 KFIAEKTGLQIAVYSKKIAQFADADKLPDVGNIQFNLIDDGKVVLLPPDAVGHTWYGT 300
 +IA+ G+ I + + D G + +F DG + L+P +G+T +GTT
 Sbjct: 236 NYIADNFGVSIVLENGTYRN-----DKGEVSKF--YDPGHLTLIFNGPLGNTVFGTT 285

Query: 301 PEAFLASGGT-DAQVQLSGGPTVTTYLEKHPVNIATVVSAMVIPSFEGIDYVGVLT 357
 PE DL + T +A+V+++ G VTT PVN+ T VS V +PSFE +D V +LT
 Sbjct: 286 PEESDLFADNTVNAEIVDNGIAVTTTCTDPVNVQTKVSMVALPSFERLDDVYMLT 343

Query= sid|114834|lan|dp1ORF013 Phage dp1 ORF|10215-11240|3
 (341 letters)

>sp|P09122|DP3X_BACSU DNA POLYMERASE III SUBUNITS GAMMA AND TAU
 Length = 563

Score = 182 bits (458), Expect = 2e-45
 Identities = 118/353 (33%), Positives = 176/353 (49%), Gaps = 31/353 (8%)

Query: 7 YRPQTFFEVVAQEYVKEILLNQLNGAIAKHGYLFCXXXXXXXXXXRIFAKDVN----- 60
 +RPQ FE+VV QE++ + L N L H YLF +IFAK VN
 Sbjct: 10 FRPQRFEDVVGQEHITKTLQNALQKKFSAHYLFSGPRGTGKTSAAKIFAKAVNCEHAPV 69

Query: 61 -----KGL-----GSPIDEAASNNGVENVRNIEDSRYKSMSEFKVYIIDEVH 105
 KG+ IEIDAASNNGV+ +R+I + ++ +KVYIIDEVH
 Sbjct: 70 DEPCNECAACKGITNGSISDVIEIDAASNNGVDEIRDIRDKVKFAPSAVTYKVYIIDEVH 129

Query: 106 MLSTGAFNALLKTLLEPSSGTVFILCTTDPQKIPDTILSRVQRFDFTRIDNDDIVNQLQF 165
 MLS GAFNALLKTLLEP +FIL TT+P KIP TI+SR QRFD RI + IV ++
 Sbjct: 130 MLSIGAFNALLKTLLEPPEHCIFILATTEPHKIPLTIISRCQRFDFKRITSQAIVGRMNK 189

Query: 166 IIESENEEGAGYSYERDALSFIGKLANGMRDSITRLEKVLVDYSHHVDMEAVSNAL---G 222
 I+++E E +L I A+GMRD+++ L++ + +S D+ V +AL G
 Sbjct: 190 IVDABEQ-----LQVEEGSLEIIASAAHGMRDALSLDQAIISFG--DILKVEDALLITG 242

Query: 223 VPDYETFASLVEAIAANYDGSCKLEIVNDFHYSKDLKLVTRNFTDFLLEVCKYWLVRDIS 282
 L +++ + + S LE +N+ GKD + + + ++ Y +
 Sbjct: 243 AVSQLYIGLAKSLHDKNVSDALETNLNELLQQGDKPAKLIEDMIFPRDMLLYKTAPGLE 302

Query: 283 ITQLPAHFESKLEQFCEAFQYPTLLWMLEEMNELAGVVKWEPNAKPIIETKLL 335
 + + E L M++ +N+ +KW + + E ++
 Sbjct: 303 GVLEKVKVDETFRELSEQIPAQALYEMIDILNKSHQEMKWTNHPRIFFEVAVV 355

Query= sid|114835|lan|dp1ORF014 Phage dp1 ORF|50961-51974|3
 (337 letters)

>sp|P47492|PRIM_MYCGE DNA PRIMASE >gi|1361496|pir||F64227 DNA primase (dnaE) homolog
 MG250 - Mycoplasma genitalium (SGC3) >gi|3844848
 (U39704) DNA primase (dnaE) [Mycoplasma genitalium]
 Length = 607

Score = 57.0 bits (135), Expect = 2e-07
 Identities = 53/190 (27%), Positives = 89/190 (45%), Gaps = 17/190 (8%)

Query: 146 EELDKYRFIHP-----YMYERKLDELIEFMFDVGYDK--LHDCITFPVRNLKGETVVF 196
 E +++Y FI+P Y++ K + + FD K + I P+ + G V F
 Sbjct: 170 ESMERYPPFINPKIKPSELYLFS-KTNQQGLGFFDFNTKKATFQNMIMPIHDFNGNPNVGF 228

Query: 197 NRRSVRSKPHQYGEDDPKTEFLYQYELVAFRDYFEKPIQVFVTSVINCLTLWSMKIP 256
 + RSV + ++ EF + + EL+ K ++Q+F+ E + TL + K
 Sbjct: 229 SARSVDNINKLKYKNSADHEF-FKKGELLFNHRLNKNLNLQLFIVEGYFDVFTLTNSKFE 287

Query: 257 AVALMGVGGGN-QINLLKR--LPYRNIVLALDPDNAGQTAQEKLYRQLKRSK-VVRFLNY 312
 AVALMG+ + QI +K + +VLALD D +GQ A L +L + +V + +
 Sbjct: 288 AVALMGLALNDVQIKAIKAHFKEQLTLVLALDNDASGQNAVFSLEKLNNNNFIVEIVQW 347

405

Query: 313 PKEFYDNKWD 322
 + D WD
 Sbjct: 348 EHNYKD--WD 355

Query= sid|114837|lan|dp1ORF016 Phage dp1 ORF|43413-44303|3
 (296 letters)

>emb|CAB07986| (Z93946) N-acetylmuramoyl-L-alanine amidase [bacteriophage Dp-1]
 Length = 296

Score = 661 bits (1686), Expect = 0.0
 Identities = 296/296 (100%), Positives = 296/296 (100%)

Query: 1 MGVDIEKGVAMQARKGRVSYSDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH 60
 MGVDIEKGVAMQARKGRVSYSDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH
 Sbjct: 1 MGVDIEKGVAMQARKGRVSYSDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH 60

Query: 61 AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMPIDSDNIIHCNYAYDGIS 120
 AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMPIDSDNIIHCNYAYDGIS
 Sbjct: 61 AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMPIDSDNIIHCNYAYDGIS 120

Query: 121 VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE 180
 VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE
 Sbjct: 121 VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE 180

Query: 181 ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYFNRDGSMTGW 240
 ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYFNRDGSMTGW
 Sbjct: 181 ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYFNRDGSMTGW 240

Query: 241 IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV 296
 IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV
 Sbjct: 241 IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV 296

Query= sid|114841|lan|dp1ORF020 Phage dp1 ORF|1864-2658|1
 (264 letters)

>emb|CAB13247| (Z99111) similar to coenzyme PQQ synthesis [Bacillus subtilis]
 Length = 243

Score = 217 bits (548), Expect = 5e-56
 Identities = 117/248 (47%), Positives = 163/248 (65%), Gaps = 15/248 (6%)

Query: 23 MPIMEIFGPTIQEGMVIGQKTIFIRTGGCDYHCNWCDSAFTWNGTTEPE--YITGKEAA 80
 +P++EIFGPTIQEGMVIGQKT+P+RT GCDY C+WCDSAFTW+G+ + + ++T +E
 Sbjct: 5 IPVLEIFGPTIQEGMVIGQKTMFVRTAGCDYSCSWCDSAFTWDGSAKKDIRWMTAEEIF 64

Query: 81 SRILKLAFNKGEQICNVTLTGGNPALINEPMAKMISILKEHGFKFGLTQGRFQEW 140
 + + D G +HVT++GGNPAL+ + + I +LKE+ + LETQGT +Q+WF
 Sbjct: 65 AEL-----KDIGDAPSHVTISGPNALLKQ-LDAFIELLKENNIRAALETQGT VYQDWF 118

Query: 141 KEVSDITISPKPPSSGMRTNMKILEAIVDRM--NDENLDWSPKIVIFDENDLAYARDMFK 198
 + D+TISPKPPSS M TN + L+ I+ + ND S K+VIF++ DL +A+ + K
 Sbjct: 119 TLIDDLTISPKPPSSKMVTNFQKLDHILTSQENDRQHAVSLKVVIFNDEDELEFAKTVHK 178

Query: 199 TFEGKLRPVNYLSVGNANAY--EEGKISDRLLEKLGWLWDKVYEDPAFNNVRPLPQLHTL 256
 + G YL VGN + + + + LL K L DKV D N VR LPQLHTL
 Sbjct: 179 RYPG---IPFYLVGNDDVHTTDDQSLIAHLLGKYEALVDKVAVDALNLVRVLPQLHTL 235

Query: 257 VYDNKRGV 264
 ++ NKRGV
 Sbjct: 236 LWGNKRGV 243

406

Query= sid|114842|lan|dp1ORF021 Phage dpl ORF|2504-3295|2
(263 letters)

>sp|P19465|GCH1_BACSU GTP CYCLOHYDROLASE I (GTP-CH-I) >gi|98411|pir||A38256 GTP
cyclohydrolase I (EC 3.5.4.16) - Bacillus subtilis
>gi|143231 (M37320) regulatory protein [Bacillus
subtilis] >gi|143799 (M80245) MtrA [Bacillus subtilis]
>gi|2634696|emb|CAB14194| (Z99115) GTP cyclohydrolase I
[Bacillus subtilis]
Length = 190

Score = 208 bits (523), Expect = 4e-53
Identities = 103/185 (55%), Positives = 133/185 (71%), Gaps = 1/185 (0%)

Query: 80 VTLDNTEAAVQRLFLGLGEDAERDGLQDTPFRFVKALAEHTVGYREDPKLHLEKTFDVDH 139
V + E AV+++ +GED R+GL DTP R K AE G EDPK H + F +H
Sbjct: 4 VNKEQIEQAVRQILEAIGEDPNREGLLDTPKRVAKMYAEVFSGLNEDPKHEFQTIIFGENH 63

Query: 140 EDLVLVKDIPIFNSLCEHHLAPFVGKVHAIYIPKD-KITGLSKFGRVVEGYAKRLQVQERL 198
E+LVLVKDI F+S+CEHHL PF GK H+AYIP+ K+TGLSK R VE AKR Q+QER+
Sbjct: 64 EELVLVKDIAFHSMEHHLVPFYGKAHVAYIPRGKVTGLSKLARAVEAVAKRPQLQERI 123

Query: 199 TQIADAIQEVLNPQAVAVIVEAEHTCMSGRGIKKGATTVTSTMRLGFQDDASARAELL 258
T IA++I E L+P V V+VEAEH CM+ RG++K GA TVTS +RG+F+DDA+ARAE+L
Sbjct: 124 TSTIAESIVETLDPHGMVVVVEAEHMCMTMRGVKPGAKTVTSVAVRGVFKDDAAARAELV 183

Query: 259 QLIKK 263
+ IK+
Sbjct: 184 EHIKR 188

Query= sid|114843|lan|dp1ORF022 Phage dpl ORF|30896-31675|2
(259 letters)

>gi|2347102 (U77367) internalin [Listeria monocytogenes]
Length = 821

Score = 55.0 bits (130), Expect = 5e-07
Identities = 44/149 (29%), Positives = 63/149 (41%), Gaps = 13/149 (8%)

Query: 119 FRMNIYVVPNYVG--DSIVNYVKITLNNCTGKAPGLSIGKEFYAPEFNIKAREATKAGLPV 176
F + VPN + D + + NN T AP L Y PE +K + K +
Sbjct: 383 FSKTSLVPMNITSIDGTIAPETISNNGTYDAPNLKWSLPNYLPE--VKYTFSQKIPIGT 440

Query: 177 KSM DYVAQLPAVLR-----RVTFDLNGGTGTADAVRVEAGKKISPKPVDPTLTGKAFKGW 231
+ +Y + L+ +VTF++ G T + V E + P+P PT G F GW
Sbjct: 441 GTSNYSGFITQPLKELLDYKVTFNVEGNTSEVETVTEE---NLIEPTSPKQGYTFDGV 497

Query: 232 -KVEGESTIWDNFDNHMPDRDVKLVAQFA 259
E T WDF MP D+ L A F+
Sbjct: 498 YDAETGGTKWDFTTGQMPANDLTLYAHFS 526

Score = 43.4 bits (100), Expect = 0.002
Identities = 47/195 (24%), Positives = 73/195 (37%), Gaps = 12/195 (6%)

Query: 72 YDLTFKDNITFDPEIMALIEGGTVRQGGGTIAGYDT-PMLAQGASNMKPFMRNIYVVPNY-- 128
YD + T + +G + GG + T MA + F +N Y N+
Sbjct: 547 YDALLNEPTTPTKQGYTFDGYDAETGGNKWDFKTMKMPANDVAFYAHFTINNYQANFDI 606

Query: 129 ---VGDSIVNYVKITLNNCTGKAPGLSIGKEFYAPEFNIKAREATKAGLPVKSM DYVAQL 185
V + + Y + T G + + A K TK +P + A
Sbjct: 607 DGEVKNETIAYDTLLNEPTTPTKQGYTFDGYDAETGGTKWDFKTKE-MPANDVTLYAHF 665

Query: 186 PAVLRRVTFDLNGGTGTADAVRVEAGKKISPKPVDPTLTGKAFKGW-KVEGESTIWDNFDN 244
+ FD++G T + V +A + P+P P+ TG +GW E T WDF
Sbjct: 666 TINNYQANFDIDGAV-TEEVVNYDA---LIPEPTSPSKTGFTLEGWYDAEVGGTKWDFKT 721

Query: 245 HMMPDRDVKLVAQFA 259
MP D+ L A F+
Sbjct: 722 MKMPANDITLYAHFS 736

Score = 38.3 bits (87), Expect = 0.057
Identities = 42/169 (24%), Positives = 59/169 (34%), Gaps = 10/169 (5%)

407

Query: 96 QQGGTIAGYDT-PMLAQGASNMKPFPMNIYVPNYVGDIVNYVKIT----LNNCTGKAPG 150
 + GGT + T M A + F +N Y N+ D +V + LN T
 Sbjct: 501 ETGGTKWDFTTGQMPANDLTLYAHFSVNSYQANFDIDGVVINEAVVYDALLNEPTTPTKQ 560

Query: 151 LSIGKEFYAPEFNKAREATKAGLPVKSMYVQALPAVLRRVTFDLNGGTGTADAVRVEA 210
 +Y E + +P + + A + FD++G A
 Sbjct: 561 GYTFDGWYDASTGGNKWDFKTMKNPANDVAFYAHFTINNYQANFDIDGEVKNETI----A 616

Query: 211 GKKISPFPVDPILTCKAFKGW-KVEGESTIWDNFMMPDRDVKLVQAF 258
 + +P PT G F GW E T WDF MP DV L A F
 Sbjct: 617 YDTLLNEPTTPTKQGYTFDGWYDAETGGTKWDFKTKEMPANDVTLYAHF 665

Query= sid|114850|lan|dp1ORF029 Phage dpl ORF|662-1348|2
 (228 letters)

>gi|2650185 (AE001074) succinoglycan biosynthesis regulator (exsB)
 [Archaeoglobus fulgidus]
 Length = 239

Score = 119 bits (295), Expect = 2e-26
 Identities = 79/224 (35%), Positives = 113/224 (50%), Gaps = 11/224 (4%)

Query: 1 MKSVLLSGGVDSATCLAIEVDKNGSKNVHAIAPNYGQKHEAELENAANVAMFYGVKFTI 60
 MK+V+LLSGG+DS+T L +D G VHA+ F YGQKH E+E+A VA V+
 Sbjct: 1 MKAVMLLSGGIDSSTLLYYLLD--GGYEVHALTFYQKHSKEIESAEKVAKAQVRHLK 58

Query: 61 LEIDSKIYXXXXXXLLQKGEISHGKSYAEILAEKEVVDYVPPFRNGLMSQXXXXXXXX 120
 ++I S I+ L G+ E+ Y+E + + T VP RN ++LS
 Sbjct: 59 VDI-STIHDLISYGALTGEBEVPA-KFYSEEVQRR---TIVPNRNMILLS--IAAGYAV 110

Query: 121 XXXXXXXXXXXXXXXXXXXXPDCTPEFYNSMSNAMEYGT-GGKVTLVAPLLTLTKAQVVK 179
 PDC EF ++ A+ V + AP + +TKA +V+
 Sbjct: 111 KIGAKEVHYAAHLSDSIYPDCRKEFVKALDTAVYLANIWTPEVRAPFVDMTKADIVRL 170

Query: 180 GIDLDPYFLTRSCYEDAESCGTCATCIDRKKAFEENGMDPI 223
 G+ L VPY LT SCYE C +C TC++R +AF NG+ DP+
 Sbjct: 171 GLKLGVPYELTWSCYEGGDRPCLSCGTCLERTEAFLANGVKDPL 214

Query= sid|114855|lan|dp1ORF034 Phage dpl ORF|131-652|2
 (173 letters)

>emb|CAB13248| (Z99111) similar to hypothetical proteins [Bacillus subtilis]
 Length = 165

Score = 220 bits (556), Expect = 4e-57
 Identities = 103/139 (74%), Positives = 117/139 (84%)

Query: 5 TTRTDABLTGVTLLGNQDTKYDYDYNPDVLETFPNKHPENNYLVTFDGYEFTSLCPKTGQ 64
 TTR ++EL GVTLLGNQ T Y ++Y PDVLE+FPNKH +Y V F+ EFTSLCPKTGQ
 Sbjct: 2 TTRKESELEGVTLLGNQGTNYLFEYAPDVLESFPNKHVNRDYFVKFNCPEFTSLCPKTGQ 61

Query: 65 PDFANVPISYIPNEKMVESKSLKLYLFSFRNHGDFHEDCMNII+NDLYELMEPKYIEVMG 124
 PDFA ++ISYIP+EKMVESKSLKLYLFSFRNHGDFHEDCMNII+NDL ELM+P+YIEV G
 Sbjct: 62 PDFATIYISYIPDEKMVESKSLKLYLFSFRNHGDFHEDCMNII+NDLIELMDPRYIEVMG 121

Query: 125 LFTPRGGISYIPFVNKVP 143
 FTTPRGGISI P+ N P
 Sbjct: 122 KFTPRGGISIDPYTNYGKP 140

Query= sid|114857|lan|dp1ORF036 Phage dpl ORF|48808-49362|1
 (184 letters)

>gi|1353529 (U38906) ORF12 [Bacteriophage rlt]
 Length = 296

Score = 53.5 bits (126), Expect = 1e-06
 Identities = 42/149 (28%), Positives = 70/149 (46%), Gaps = 9/149 (6%)

Query: 34 IASNTVGNKGTSWAVRLLQRYLAETALDGRIVEKGMFVVSAQLLTFEGDYNFYQTMQEF 93
 + S G GK+ A+ +L+ L T L ++ V + F + + F + + F+
 Sbjct: 155 VVSGPAGTGKSHLAMSILKDCLOHTDLT--VIFASWSEVLHLIKDSFDNKSDFYSTYEFM 212

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Query: 94 ERFERLKTCELLVIDEIGGSLTKASYFYLYDLVNYRVDNNLSTIYTTNYTDDEIIDLLG 153
 E F + +LLVID+IG +T+ S L ++++ R TI TTN DEI
 Sbjct: 213 EVF---RNTDLLVIDDIGSEKITEWSMSLLTEVLDART----KTIITTNLKSDEIRKKYH 265

Query: 154 QRLYSRIYDTSVVLDFOASNVRLGVSEI 182
 R YSR++ F N++ VS++
 Sbjct: 266 NRTYSRLFRGIGKKAFNFENIKDKRVSQL 294

Query= sid|114859|lan|dp1ORF038 Phage dp1 ORF|1350-1871|3
 (173 letters)

>sp|P44123|YB90_HAEIN_HYPOTHETICAL_PROTEIN_HI1190 >gi|1074675|pir||F64021 hypothetical
 protein HI1190 - Haemophilus influenzae (strain Rd KW20)
 >gi|1574117 (U32798) 6-pyruvoyl tetrahydrobiopterin
 synthase, putative [Haemophilus influenzae Rd]
 Length = 141

Score = 100 bits (247), Expect = 6e-21
 Identities = 59/143 (41%), Positives = 83/143 (57%), Gaps = 10/143 (6%)

Query: 2 RVSKTLTFDAAHQVLGHFGKCANLHGHTYKVEISLAGGTYDHGSSQGMVVDYFHVKKIA- 60
 ++SK +FD AH L GH GKC NLHGHTYK+++ ++G Y G+ + MV+DF +K I
 Sbjct: 3 KISKEFSFDMHLLDGHGDKCQNLHGHTYKQVEISGDLVKSGAKKAMVIDFSDLKSIVK 62

Query: 61 GTFIDRLDHAVALL-QGNEP----IALANAVDTKRVLFGRFRTTAENMSRFLTWLTLMWK 115
 +D +DHA + Q NE L +++K FRITAE ++RF+ L +
 Sbjct: 63 KVILPMDHAFIYDQTNERESQIATLLQKLNSKTFGVFPRTTAEIARFIFNRLKH--DE 120

Query: 116 HARIDSIKLWETPTGCAECTYEE 138
 I SI+LWETPT + C Y E
 Sbjct: 121 QLSISSIRLWETPT--SFCEYQE 141

Query= sid|114860|lan|dp1ORF039 Phage dp1 ORF|3306-3803|3
 (165 letters)

>emb|CAA68244| (X99978) ORF7; hydrophobic protein [Lactobacillus plantarum]
 Length = 168

Score = 64.4 bits (154), Expect = 5e-10
 Identities = 49/156 (31%), Positives = 84/156 (53%), Gaps = 9/156 (5%)

Query: 8 WLVRTALIAALYVTLTVAFSAISY--GPIQFRVSEALILLPLMNHRTWPGIVLGTIIANF 65
 W++ AL+AA+YV L + +A S G IQFRVSE L L ++N ++ GIV G I+ +
 Sbjct: 9 WIIN-ALVAAMYVVLCLGPAAFSLASGAIQFRVSEGLNHLAVFNRYKINGIVAGVILFDA 67

Query: 66 FSP-LGLIDVLFGSLATFLGXXXXXXXXXXSPLYSLICPVLA----NAYLIALELRIVY 120
 F P L++VLF + L ++ + +A + ++IAL + ++
 Sbjct: 68 FGPAGSLNVLFGGQSLLALLVLTWLAPKLKTWQRMLLNIALFTVSMFMIALMITMS 127

Query: 121 S-LPFWESVIYVGISEAIIVLISYFLISTLAKNNHF 155
 S + FW + + +SE II+ I+ ++ +L + HF
 Sbjct: 128 SGVAFWPPTYLTALSELIIMSITAPIMYSLDRVLHF 163

Query= sid|114862|lan|dp1ORF041 Phage dp1 ORF|8208-8699|3
 (163 letters)

>gi|2522313 (AF012906) dUTPase homolog [Bacillus subtilis]
 >gi|2634394|emb|CAB13893| (Z99114) similar to
 deoxyuridine 5'-triphosphate nucleotidohydrolase
 [Bacillus subtilis] >gi|3025643 (AF020713) putative
 dUTPase [Bacteriophage SPBc2]
 Length = 142

Score = 108 bits (267), Expect = 2e-23
 Identities = 65/160 (40%), Positives = 83/160 (51%), Gaps = 25/160 (15%)

Query: 5 VDVKMIDPKLDRLKYT--GDWVDVRISITKIDADSADVSRCKVLQKAQVYSVAAGECI 62
 + +K +D R+ GDW+D+R + I D +
 Sbjct: 3 IKIKYLDQTRINKMEQGDWIDLRAEDVAIKKDEFL----- 41

Query: 63 KIAHGFALELPKGYEAILHPRSSLFKKTGLIFVSS-GVIDEGYKGDTEWFSVWYATRDA 121
 + G A+ELP+GYEA + PRSS +K G+I +S GVIDE YKGD D WF YA RD
 Sbjct: 42 -VPLGVAMELPEGYEAHVVPSSSTYKNFGVIQTNSMGVIDESYKGDNDFWFFPAYALRDT 100

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Query: 122 DIFYDQRIAQFRIQEKQPAIKFNFVESLGNAARGGHGSG 161
I RI QFRI +K PA+ V+ LGN RGGHSGSG
Sbjct: 101 KIKKGDRICQFRIMKKMPAVDLIEVDRLGNGDRGGHSGSG 140

Query= sid|114867|lan|dp1ORF046 Phage dp1 ORF|42774-43202|3
(142 letters)

>emb|CAB07984| (Z93946) hypothetical protein [bacteriophage Dp-1]
Length = 142

Score = 287 bits (728), Expect = 2e-77
Identities = 142/142 (100%), Positives = 142/142 (100%)

Query: 1 MPMWLNDTAVLTTIITACSGVLTVLLNKLFEWKSNAKSVLEDISTTLSTLKQQVDGIDQ 60
MPMWLNDTAVLTTIITACSGVLTVLLNKLFEWKSNAKSVLEDISTTLSTLKQQVDGIDQ
Sbjct: 1 MPMWLNDTAVLTTIITACSGVLTVLLNKLFEWKSNAKSVLEDISTTLSTLKQQVDGIDQ 60
Query: 61 TTVAINHQNQNDVIQDGTTRKIQRVRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE 120
TTVAINHQNQNDVIQDGTTRKIQRVRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE
Sbjct: 61 TTVAINHQNQNDVIQDGTTRKIQRVRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE 120
Query: 121 VEALYEKYKKLPIREEDLDETI 142
VEALYEKYKKLPIREEDLDETI
Sbjct: 121 VEALYEKYKKLPIREEDLDETI 142

Query= sid|114901|lan|dp1ORF080 Phage dp1 ORF|42490-42759|1
(89 letters)

>emb|CAB07983| (Z93946) hypothetical protein [bacteriophage Dp-1]
Length = 124

Score = 147 bits (367), Expect = 1e-35
Identities = 75/75 (100%), Positives = 75/75 (100%)

Query: 1 MLNLTCSRQIVAEFTIGQGAEEKLVKTTIVNIDANAVSTVSETLHDPDLAANRRELAD 60
MLNLTCSRQIVAEFTIGQGAEEKLVKTTIVNIDANAVSTVSETLHDPDLAANRRELAD
Sbjct: 1 MLNLTCSRQIVAEFTIGQGAEEKLVKTTIVNIDANAVSTVSETLHDPDLAANRRELAD 60
Query: 61 EQKLRETRYAIEDEI 75
EQKLRETRYAIEDEI
Sbjct: 61 EQKLRETRYAIEDEI 75

Query= sid|114912|lan|dp1ORF091 Phage dp1 ORF|43189-43413|1
(74 letters)

>emb|CAB07985| (Z93946) holin [bacteriophage Dp-1]
Length = 74

Score = 63.2 bits (151), Expect = 2e-10
Identities = 34/74 (45%), Positives = 34/74 (45%)

Query: 1 MKLSNEQYDXXXXXXXXXXXXXXXXXXXXXXXXXXQFDXXXXXXXXXXXXXXXXXXVLGVSSR 60
MKLSNEQYD YQFD VLGVSRR
Sbjct: 1 MKLSNEQYDVAKNVVTVVVPAALITGLGALYQFDTTATGTIALLATFAGTVLGVSRR 60
Query: 61 NYQKEQEAQNNEVE 74
NYQKEQEAQNNEVE
Sbjct: 61 NYQKEQEAQNNEVE 74

Condensed listing of homology information from above

Phage: dpl

Database: nr

Program: Blastp

Query= sid|114822|lan|dplORF001 Phage dpl ORF|36698-40390|2
(1230 letters)

gi 2444124	(U88974) ORF45 [Streptococcus thermophilus temperate ...	467	e-130
gi 928828	(L44593) ORF1904; putative [Lactococcus lactis phage B...	427	e-118
gi 2935676	(AF032121) unknown [Streptococcus thermophilus bacter...	309	1e-82
gi 2935691	(AF032122) unknown [Streptococcus thermophilus bacter...	306	7e-82
gi 3540289	(AF057033) putative anti-receptor [Streptococcus ther...	279	6e-74
gi 4530154	[gb AAD21894.1 (AF085222) putative tail-host specific...	220	3e-56
gi 930045	[emb CAA33387 (X15332) alpha-1 (III) collagen [Homo sa...	58	4e-07
gi 1070603	[pir CGHU7L collagen alpha 1(III) chain precursor - h...	58	4e-07
gi 4502951	[ref NP_000081.1 PCOL3A1 collagen, type III, alpha 1 ...	58	4e-07
gi 115290	[sp P04258 CA13_BOVIN COLLAGEN ALPHA 1(III) CHAIN >gi 7...	58	4e-07
gi 575322	[emb CAA36279 (X52046) type III collagen [Mus musculus]	57	8e-07
gi 2119163	[pir S59856 collagen alpha 1(III) chain precursor - m...	57	8e-07
gi 543912	[sp P13941 CA13_RAT COLLAGEN ALPHA 1(III) CHAIN >gi 543...	57	1e-06
gi 3171998	[emb CAA06510 (AJ005395) collagen alpha 1 (III) [Ratt...	57	1e-06
gi 3947565	[emb CAA90250 (Z49967) similar to collagen; cDNA EST ...	54	7e-06
gi 423403	[pir A46053 bullous pemphigoid antigen, BPAG2, type XV...	53	9e-06
gi 115410	[sp P12114 CCS1_CAEEL CUTICLE COLLAGEN SQT-1 >gi 84437 ...	53	9e-06
gi 3873801	[emb CAA90084 (Z49907) cuticle collagen SQT-1; cDNA E...	53	9e-06

Query= sid|114823|lan|dplORF002 Phage dpl ORF|32386-35835|1
(1149 letters)

gi 3341922	[dbj BAA31888 (AB009866) orf 15 [bacteriophage phi PVL]	280	3e-74
gi 4126622	[dbj BAA36642.1 (AB016282) ORF36 [bacteriophage phi-105]	232	1e-59
gi 1369948	[emb CAA59194 (X84706) host interacting protein [Bact...	201	3e-50
gi 3139112	(AF063097) gpt [Bacteriophage P2]	188	2e-46
gi 3337272	(U32222) G protein [Bacteriophage 186]	161	3e-38
gi 4063799	[dbj BAA36253 (AB008550) orf25; similar to T gene of ...	159	8e-38
gi 3172274	(AF022214) minor tail subunit; putative tape-measure ...	123	6e-27
gi 465127	[sp Q05233 VG26_BPML5 MINOR TAIL PROTEIN GP26 >gi 41904...	108	2e-22
gi 3540284	(AF057033) putative minor tail protein [Streptococcus...	99	2e-19
gi 2444119	(U88974) ORF40 [Streptococcus thermophilus temperate ...	90	6e-17
gi 2634555	[emb CAB14053 (Z99115) yomI [Bacillus subtilis] >gi 3...	66	1e-09
gi 2392838	(AF011378) unknown [Bacteriophage sk1]	64	5e-09
gi 2764873	[emb CAA66557 (X97918) gene 18.1 [Bacteriophage SPP1]	62	3e-08
gi 1353559	(U38906) ORF42 [Bacteriophage rlt]	61	6e-08
gi 630841	[pir S39079 puff C-8 protein - fungus gnat (Rhynchosci...	55	2e-06
gi 1730865	[sp P51731 YO27_BPHP1 HYPOTHETICAL 72.8 KD PROTEIN IN ...	53	8e-06
gi 224288	[prf 1101273J ORF 7 [Bacteriophage HP1]	53	1e-05

Query= sid|114824|lan|dplORF003 Phage dpl ORF|53538-55877|3
(779 letters)

gi 118825	[sp P00582 DPO1_ECOLI DNA POLYMERASE I (POL I) >gi 6705...	193	3e-48
gi 2982102	[pdb 1KFS A Chain A, All-Oxygen Dna Complexed To The 3...	193	3e-48
gi 229889	[pdb 1DPI DNA Polymerase I (Klenow Fragment) (E.C.2....	193	3e-48
gi 1169402	[sp P43741 DPO1_HAEIN DNA POLYMERASE I (POL I) >gi 107...	191	1e-47
gi 2688462	(AE001156) DNA polymerase I (polA) [Borrelia burgdorf...	190	3e-47
gi 809180	[pdb 1KLN A Escherichia coli	190	3e-47
gi 1913934	[emb CAA72997 (Y12328) DNA-directed DNA polymerase I ...	189	8e-47
gi 4090935	(AF028719) DNA polymerase type I [Rhodothermus sp. 'I...	175	1e-42
gi 4731571	[gb AAD2805.1 AF121780_1 (AF121780) DNA polymerase I ...	174	2e-42
gi 1633576	(U57757) similar to proofreading 3'-5' exonuclease an...	173	4e-42
gi 3322368	(AE001195) DNA polymerase I (polA) [Treponema pallidum]	172	9e-42
gi 1006595	[dbj BAA10748 (D64005) DNA polymerase I [Synecocysti...	171	2e-41
gi 585062	[sp Q07700 DPO1_MYCTU DNA POLYMERASE I (POL I) >gi 4161...	163	5e-39
gi 4376908	[gb AAD18751 (AE001645) DNA Polymerase I [Chlamydia p...	157	2e-37
gi 1169403	[sp P46835 DPO1_MYCLE DNA POLYMERASE I (POL I) >gi 107...	152	7e-36
gi 2145839	[pir S72949 DNA polymerase I - Mycobacterium leprae >...	152	7e-36
gi 1405438	[emb CAA67184 (X98575) DNA-dependent DNA polymerase [...	152	9e-36
gi 2506365	[sp P80194 DPO1_THECA DNA POLYMERASE I, THERMOSTABLE (...	147	2e-34
gi 3328929	(AE001322) DNA Polymerase I [Chlamydia trachomatis]	147	3e-34

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gi 3913510 sp O52225 DPO1_THEFI DNA POLYMERASE I, THERMOSTABLE (...)	146	7e-34
gi 1205984 U33536 DNA polymerase I [Bacillus stearothermophilus]	146	7e-34
gi 118827 sp P13252 DPO1_STRPN DNA POLYMERASE I (POL I) >gi 9802...	145	9e-34
gi 1942202 pdb 1JXE Stoffel Fragment Of Taq Dna Polymerase I	145	1e-33
gi 1943520 pdb 1KTQ Dna Polymerase	145	1e-33
gi 1084022 pir JX0359 DNA-directed DNA polymerase (EC 2.7.7.7) ...	145	1e-33
gi 507891 dbj BAA06775 (D32013) DNA Polymerase [Thermus aquaticus]	145	1e-33
gi 118828 sp P19821 DPO1_THEAQ DNA POLYMERASE I, THERMOSTABLE (T...	145	1e-33
gi 1706502 sp P52028 DPO1_THETH DNA POLYMERASE I, THERMOSTABLE (...)	144	2e-33
gi 1097211 prf 2113329A DNA polymerase [Thermus aquaticus therm...	144	2e-33
gi 2098289 pdb 1TAU A Chain A, Structure Of Dna Polymerase	143	3e-33

Query= sid|114825|lan|dp1ORF004 Phage dp1 ORF|40401-42440|3
(679 letters)

gi 1934761 emb CAB07981 (Z93946) hypothetical protein [bacterio...	1011	0.0
gi 3540290 AF057033 putative minor structural protein [Strepto...	346	2e-94
gi 2444125 U88974 ORF46 [Streptococcus thermophilus temperate ...]	339	3e-92
gi 1934762 emb CAB07982 (Z93946) hypothetical protein [bacterio...	300	2e-80
gi 4530155 gb AAD21895.1 (AF085222) unknown [Streptococcus ther...	276	4e-73
gi 2935677 AF032121 unknown [Streptococcus thermophilus bacter...	250	3e-65
gi 2935692 AF032122 unknown [Streptococcus thermophilus bacter...	250	3e-65
gi 1136289 U42597 histidine kinase A [Dictyostelium discoideum]	50	7e-05

Query= sid|114827|lan|dp1ORF006 Phage dp1 ORF|45296-46987|2
(563 letters)

gi 4377165 gb AAD18987 (AE001666) SWI/SNF family helicase_2 [Ch...	171	1e-41
gi 1769947 emb CAA67095 (X98455) SNF [Bacillus cereus]	160	3e-38
gi 3329163 AE001341 SWF/SNF family helicase [Chlamydia trachom...	159	6e-38
gi 4377149 gb AAD18973 (AE001664) SWI/SNF family helicase_1 [Ch...	157	2e-37
gi 3328995 AE001326 SWI/SNF family helicase [Chlamydia trachom...	153	2e-36
gi 2493354 sp P75093 Y018 MYCPN HYPOTHETICAL HELICASE MG018/MG01...	146	4e-34
gi 1653748 dbj BAA18659 (D90916) helicase of the snf2/rad54 fam...	143	3e-33
gi 1763712 emb CAB05939 (Z83337) member of the SNF2 helicase fa...	143	4e-33
gi 2636153 emb CAB15645.1 (Z99122) similar to SNF2 helicase [Ba...	143	4e-33
gi 2909552 emb CAA17284 (AL021924) helz [Mycobacterium tubercul...	140	2e-32
gi 3844627 U39681 ATP-dependent RNA helicase, putative [Mycopl...	136	3e-31
gi 1351463 sp P47264 Y018 MYCGE HYPOTHETICAL HELICASE MG018	136	4e-31
gi 2660669 AC002342 human Mi-2 autoantigen-like protein [Arabi...	131	2e-29
gi 1361537 pir I64201 helicase (mot1) homolog - Mycoplasma geni...	129	4e-29
gi 3482977 emb CAA20533.1 (AL031369) putative protein [Arabidop...	128	9e-29
gi 3298562 U91543 zinc-finger helicase [Homo sapiens]	120	2e-26
gi 3875971 emb CAB02491 (Z80344) similar to helicase; cDNA EST ...	120	2e-26
gi 4557451 ref NP_001263.1 PCHD3 chromodomain helicase DNA bind...	120	2e-26
gi 2645435 AF007780 CHD3 [Drosophila melanogaster]	118	1e-25
gi 3875165 emb CAA91798 (Z67881) Similarity to Mouse Chromodoma...	118	1e-25

Query= sid|114828|lan|dp1ORF007 Phage dp1 ORF|22230-23621|3
(463 letters)

gi 2444105 U88974 ORF26 [Streptococcus thermophilus temperate ...]	89	7e-17
gi 3318666 U19754 BBA31 homolog [Borrelia burgdorferi]	59	7e-08
gi 2690260 AE000790 conserved hypothetical protein [Borrelia b...	56	5e-07

Query= sid|114829|lan|dp1ORF008 Phage dp1 ORF|49624-50961|1
(445 letters)

gi 4406210 gb AAD19901 (AF100420) DnaB replication fork helicase...	68	2e-10
gi 3121983 sp O25916 DNAB_HELFPY REPLICATIVE DNA HELICASE >gi 231...	67	2e-10
gi 4416322 gb AAD20314 (AF106032) replicative helicase; DnaB [B...	65	9e-10
gi 4155895 AE001551 REPLICATIVE DNA HELICASE [Helicobacter pyl...	60	4e-08
gi 3322317 AE001191 replicative DNA helicase (dnaB) [Treponema...	58	1e-07
gi 138031 sp P04530 VG41_BPT4 PRIMASE-HELICASE (PROTEIN GP41) >g...	53	3e-06
gi 2983861 AE000742 replicative DNA helicase [Aquifex aeolicus]	51	1e-05

Query= sid|114831|lan|dp1ORF010 Phage dp1 ORF|8699-9859|2
(386 letters)

gi 2760912 AF037258 RecA protein [Chlorobium tepidum]	133	2e-30
gi 3219851 sp P94666 RECA_CLOPE RECA PROTEIN >gi 1698591 (U61497...	129	3e-29
gi 1350566 sp P48295 RECA_STRVL RECA PROTEIN >gi 508860 (U04837)...	128	7e-29
gi 744163 prf 2014250A reca-like protein [Streptomyces violaceus]	126	3e-28
gi 730487 sp P41054 RECA_STRAM RECA PROTEIN >gi 511133 emb CAA82...	125	4e-28
gi 2687334 emb CAA15875 (AL020958) RecA protein [Streptomyces c...	125	6e-28
gi 1350565 sp P48294 RECA_STRLI RECA PROTEIN >gi 481482 pir S38...	125	6e-28

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gi|464599|sp|P33542|RECA_AQUFY RECA PROTEIN >gi|1086167|pir||A55... 123 2e-27
 gi|417636|sp|P32725|RECA_RHOSH RECA PROTEIN >gi|541307|pir||S415... 123 2e-27
 gi|2984348|AE000775| recombination protein RecA [Aquifex aeolicus] 123 2e-27
 gi|3219854|sp|P95846|RECA_STRRM RECA PROTEIN >gi|1729800|emb|CAA... 122 4e-27
 gi|2500086|sp|Q59560|RECA_MYCSM RECA PROTEIN >gi|1430892|emb|CAA... 122 4e-27
 gi|1350567|sp|P48296|RECA_THEAQ RECA PROTEIN >gi|1072963|pir||A5... 122 6e-27
 gi|625663|pir||JX0292| recA protein - Thermus aquaticus (strain HB8) 121 1e-26
 gi|1172880|sp|P42440|RECA_CAMJE RECA PROTEIN >gi|2119991|pir||I4... 120 2e-26
 gi|4154654|AE001453| RECA PROTEIN. [Helicobacter pylori J99] 120 2e-26
 gi|1072968|pir||C55020| recA protein - Thermus sp >gi|458472|dbj|... 120 2e-26
 gi|3219852|sp|P95469|RECA_PARDE RECA PROTEIN >gi|1825468|U59631... 119 3e-26
 gi|2507284|sp|P42445|RECA_HELPY RECA PROTEIN >gi|2313235|gb|AAD0... 119 4e-26
 gi|1172890|sp|Q02350|RECA_STAAU RECA PROTEIN >gi|463285|L25893|... 118 5e-26
 gi|4416209|gb|AAD20261| (AF094756) RecA protein [Bifidobacterium... 118 5e-26
 gi|2500084|sp|Q59180|RECA_BORBU RECA PROTEIN >gi|1276443|U23457... 118 5e-26

Query= sid|114832|lan|dp1ORF011 Phage dp1 ORF|28017-29096|3
 (359 letters)

gi|2444110|U88974| ORF31 [Streptococcus thermophilus temperate ... 187 1e-46
 gi|3320438|AF057033| gp348 [Streptococcus thermophilus bacterio... 179 2e-44
 gi|479514|pir||S34244| hypothetical protein p38 - actinophage VWB... 62 8e-09

Query= sid|114834|lan|dp1ORF013 Phage dp1 ORF|10215-11240|3
 (341 letters)

gi|580855|emb|CAA29958| (X06803) dnaZX-like ORF put. DNA polymer... 182 2e-45
 gi|118807|sp|P09122|DP3X_BACSU DNA POLYMERASE III SUBUNITS GAMMA... 182 2e-45
 gi|98292|pir||S13786| DNA-directed DNA polymerase (EC 2.7.7.7) II... 182 2e-45
 gi|1527142|U66040| DNA polymerase III gamma subunit [Salmonella... 172 4e-42
 gi|2494197|sp|P74876|DP3X_SALTY DNA POLYMERASE III SUBUNITS GAMM... 172 4e-42
 gi|118808|sp|P06710|DP3X_ECOLI DNA POLYMERASE III SUBUNITS GAMMA... 170 1e-41
 gi|4155207|AE001497| DNA POLYMERASE III SUBUNITS GAMMA AND TAU ... 169 2e-41
 gi|2313841|gb|AAD07767.1| (AE000584) DNA polymerase III gamma an... 168 4e-41
 gi|2583049|AF025391| DNA polymerase III holoenzyme tau subunit ... 166 3e-40
 gi|2984127|AE000759| DNA polymerase III gamma subunit [Aquifex ... 166 3e-40
 gi|3861390|emb|CAA15289| (AJ235273) DNA POLYMERASE III SUBUNITS ... 165 5e-40
 gi|1169397|sp|P43746|DP3X_HABIN DNA POLYMERASE III SUBUNITS GAMM... 156 2e-37
 gi|1293572|U49738| DNA polymerase III tau homolog DnaX [Cauloba... 151 8e-36
 gi|3328753|AE001306| DNA Pol III Gamma and Tau [Chlamydia trach... 148 4e-35
 gi|4376294|gb|AAD18193| (AE001589) DNA Polymerase III Gamma and ... 148 5e-35
 gi|581255|emb|CAA28175| (X04487) alternate dnaZX protein (AA 1-6... 146 3e-34
 gi|2688379|AE001151| DNA polymerase III, subunits gamma and tau... 140 2e-32
 gi|3323329|AE001268| DNA polymerase III, subunits gamma and tau... 137 1e-31

Query= sid|114835|lan|dp1ORF014 Phage dp1 ORF|50961-51974|3
 (337 letters)

gi|1346796|sp|P47492|PRIM_MYCGE DNA PRIMASE >gi|1361496|pir||F64... 57 2e-07
 gi|740008|prf||2004290A| primase [Haemophilus influenzae] 51 1e-05
 gi|1172619|sp|Q08346|PRIM_HABIN DNA PRIMASE >gi|1074033|pir||A64... 51 1e-05
 gi|1709769|sp|Q04505|PRIM_LACLA DNA PRIMASE >gi|1075726|pir||JC2... 51 1e-05
 gi|639846|dbj|BAA03516| (D14690) DNA primase [Lactococcus lactis] 51 1e-05

Query= sid|114837|lan|dp1ORF016 Phage dp1 ORF|43413-44303|3
 (296 letters)

gi|1934766|emb|CAB07986| (Z93946) N-acetylmuramoyl-L-alanine ami... 661 0.0
 gi|113676|sp|P06653|ALYS_STRPN AUTOLYSIS (N-ACETYLMURAMOYL-L-ALA... 221 4e-57
 gi|282326|pir||A42935| N-acetylmuramoyl-L-alanine amidase (EC 3.5... 219 3e-56
 gi|416618|sp|P32762|ALYS_BPHB3 LYTIC AMIDASE (N-ACETYLMURAMOYL-L... 212 2e-54
 gi|285273|pir||A42936| N-acetylmuramoyl-L-alanine amidase (EC 3.5... 212 2e-54
 gi|127787|sp|P15057|LYCA_BPCP1 LYSOZYME (ENDOLYSIS) (MURAMIDASE)... 162 4e-39
 gi|67761|pir||MUBPCP| N-acetylmuramoyl-L-alanine amidase (EC 3.5... 162 4e-39
 gi|127789|sp|P19386|LYCA_BPCP9 LYSOZYME (ENDOLYSIS) (MURAMIDASE)... 160 1e-38
 gi|928832|L44593| ORF259; putative [Lactococcus lactis phage BK... 119 2e-26
 gi|2511705|emb|CAA71783| (Y10818) sigA binding protein [Streptoc... 111 9e-24
 gi|4097980|U72655| surface protein C [Streptococcus pneumoniae] 107 1e-22
 gi|2351768|U89711| PspA [Streptococcus pneumoniae] 105 4e-22
 gi|2425109|AF019904| choline binding protein A [Streptococcus p... 104 6e-22
 gi|282335|pir||A41971| surface protein pspA precursor - Streptoco... 104 1e-21
 gi|2576331|emb|CAA05158| (AJ002054) SpsA protein [Streptococcus ... 103 2e-21
 gi|2127295|pir||S57962| cspC protein - Clostridium acetobutylicum... 85 6e-16
 gi|2576333|emb|CAA05159| (AJ002055) SpsA protein [Streptococcus ... 84 1e-15
 gi|4106522|gb|AAD02874.1| (AF097909) excreted protein FibB [Pept... 83 3e-15
 gi|1361406|pir||S57714| cspB protein - Clostridium acetobutylicum... 82 4e-15
 gi|1914872|emb|CAB04758| (Z82001) PCPA [Streptococcus pneumoniae] 81 9e-15

413

gi 3168594 dbj BAA28613 (AB012763) SpaA (Erysipelothrix rhusiop...	81	1e-14
gi 2292750 emb CAA64942 (X95646) homology to orf259 of lactococ...	80	3e-14
gi 2935696 (AF032122) putative lysin (Streptococcus thermophilus...	80	3e-14
gi 4586910 dbj BAA76540.1 (AB017447) protective antigen SpaA.1 ...	80	3e-14
gi 3540294 (AF057033) lysin (Streptococcus thermophilus bacterio...	79	5e-14

Query= sid|114841|lan|dp1ORF020 Phage dp1 ORF|1864-2658|1
(264 letters)

gi 2633745 emb CAB13247 (Z99111) similar to coenzyme PQQ synthe...	217	5e-56
gi 2808502 emb CAA12532 (AJ225561) ExsD protein (Sinorhizobium ...	163	1e-39
gi 3861151 emb CAA15051 (AJ235272) unknown (Rickettsia prowazekii)	82	6e-15
gi 1652793 dbj BAA17712 (D90908) hypothetical protein (Synechoc...	76	3e-13
gi 1723815 sp P55139 YGCF_ECOLI HYPOTHETICAL 25.0 KD PROTEIN IN ...	70	2e-11
gi 2984272 (AE000769) hypothetical protein (Aquifex aeolicus)	66	4e-10
gi 4155435 (AE001516) putative (Helicobacter pylori J99)	57	1e-07
gi 2127833 pir C64505 coenzyme PQQ synthesis protein III homolo...	55	5e-07
gi 2622338 (AE000890) coenzyme PQQ synthesis protein III (Methan...	54	9e-07
gi 3257042 dbj BAA29725 (AP000003) 254aa long hypothetical prot...	53	2e-06
gi 2314068 gb AAD07976.1 (AE000602) conserved hypothetical prot...	52	6e-06
gi 1723816 sp P45097 YGCF_HAEIN HYPOTHETICAL PROTEIN HI1189 >gi ...	50	2e-05

Query= sid|114842|lan|dp1ORF021 Phage dp1 ORF|2504-3295|2
(263 letters)

gi 127481 sp P19465 GCH1_BACSU GTP CYCLOHYDROLASE I (GTP-CH-I) >...	208	4e-53
gi 3242315 emb CAA04237 (AJ000685) GTP cyclohydrolase (Streptoc...	191	4e-48
gi 2494695 sp Q54769 GCH1_SYNP7 GTP CYCLOHYDROLASE I (GTP-CH-I) ...	189	2e-47
gi 255061 bbs 112832 (S44049) GTP cyclohydrolase I (clone hGCH-1...	187	7e-47
gi 4503949 ref NP_000152.1 PGCH1 GTP cyclohydrolase 1 (dopa-res...	187	7e-47
gi 2113967 emb CAB08935 (Z95557) folE (Mycobacterium tuberculosis)	187	7e-47
gi 1730240 sp P50141 GCH1_CHICK GTP CYCLOHYDROLASE I (GTP-CH-I) ...	185	3e-46
gi 2494696 sp Q55759 GCH1_SYNY3 GTP CYCLOHYDROLASE I (GTP-CH-I) ...	184	5e-46
gi 121061 sp P22288 GCH1_RAT GTP CYCLOHYDROLASE I PRECURSOR (GTP...	184	6e-46
gi 3183014 sp O13774 GCH1_SCHPO GTP CYCLOHYDROLASE I (GTP-CH-I) ...	184	6e-46
gi 3097224 emb CAA18795 (AL023093) GTP cyclohydrolase I (Mycoba...	182	2e-45
gi 2494697 sp Q19980 GCH1_CARL PROBABLE GTP CYCLOHYDROLASE I (G...	182	2e-45
gi 462167 sp Q05915 GCH1_MOUSE GTP CYCLOHYDROLASE I PRECURSOR (G...	180	7e-45
gi 1669664 emb CAA89808 (Z49706) GTP cyclohydrolase I (Dictyost...	180	1e-44
gi 2981082 (AF052048) GTP-cyclohydrolase (Ostertagia ostertagi)	178	3e-44
gi 31954 emb CAA78908 (Z16418) GTP cyclohydrolase I (Homo sapi...	177	8e-44
gi 551344 bbs 150280 (S71373) GTP cyclohydrolase I (mice, Peptid...	174	5e-43
gi 1730247 sp P51601 GCH1_YEAST GTP CYCLOHYDROLASE I (GTP-CH-I) ...	174	7e-43
gi 1246912 emb CAA87397 (Z47201) GTP cyclohydrolase 1 (Saccharo...	172	2e-42
gi 1730246 sp P51595 GCH1_STRPN GTP CYCLOHYDROLASE I (GTP-CH-I) ...	168	3e-41
gi 2982951 (AE000680) GTP cyclohydrolase I (Aquifex aeolicus)	164	6e-40

Query= sid|114843|lan|dp1ORF022 Phage dp1 ORF|30896-31675|2
(259 letters)

gi 2347102 (U77367) internalin (Listeria monocytogenes)	55	5e-07
gi 3123226 sp P25146 INLA_LISMO INTERNALIN A PRECURSOR >gi 48705...	52	4e-06
gi 149674 (M67471) internalin (Listeria monocytogenes)	52	4e-06

Query= sid|114850|lan|dp1ORF029 Phage dp1 ORF|662-1348|2
(228 letters)

gi 2650185 (AE001074) succinoglycan biosynthesis regulator (exsB...	119	2e-26
gi 3861231 emb CAA15131 (AJ235272) unknown (Rickettsia prowazekii)	117	8e-26
gi 2622210 (AE000881) conserved protein (Methanobacterium thermo...	108	4e-23
gi 2983380 (AE000709) trans-regulatory protein ExsB (Aquifex aeo...	88	6e-17
gi 1001327 dbj BAA10814 (D64006) ExsB (Synechocystis sp.)	88	6e-17
gi 2128055 pir B64468 hypothetical protein homolog MJ1347 - Met...	83	1e-15
gi 4155143 (AE001491) putative (Helicobacter pylori J99)	82	4e-15
gi 2313760 gb AAD07701.1 (AE000578) conserved hypothetical prot...	80	2e-14
gi 2120814 pir S60183 protein ExsB - Rhizobium meliloti >gi 114...	76	3e-13
gi 2633743 emb CAB13245 (Z99111) similar to hypothetical protei...	75	5e-13
gi 1175543 sp P44124 YBAX_HAEIN HYPOTHETICAL PROTEIN HI1191 >gi ...	74	1e-12
gi 2495537 sp P77756 YBAX_ECOLI HYPOTHETICAL 25.5 KD PROTEIN IN ...	71	5e-12
gi 3256471 dbj BAA29154.1 (AP000001) 269aa long hypothetical pr...	67	1e-10
gi 2921156 (AF022216) aluminum resistance protein (Arthrobacter ...	54	1e-06

Query= sid|114855|lan|dp1ORF034 Phage dp1 ORF|131-652|2
(173 letters)

gi 2633746 emb CAB13248 (Z99111) similar to hypothetical protei...	220	4e-57
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4|4

gi 4155926 (AE001554) putative (Helicobacter pylori J99]	162	1e-39
gi 2314588 gb AAD08456.1 (AE000642) conserved hypothetical prot...	161	3e-39
gi 2983458 (AE000714) hypothetical protein [Aquifex aeolicus]	103	9e-22
gi 1006604 dbj BAA10757 (D64005) hypothetical protein [Synechoc...	87	6e-17
gi 2967529 (U11045) unknown [Buchnera aphidicola]	79	2e-14
gi 2495654 sp Q46920 YQCD_ECOLI HYPOTHETICAL 32.6 KD PROTEIN IN ...	69	2e-11
gi 1175604 sp P44153 YQCD_HAEIN HYPOTHETICAL PROTEIN HI1291 >gi ...	63	1e-09
gi 3860642 emb CAA14543 (AJ235270) unknown [Rickettsia prowazekii]	56	1e-07
Query= sid 114857 lan dp1ORF036 Phage dp1 ORF 48808-49362 1 (184 letters)		
gi 1353529 (U38906) ORF12 [Bacteriophage rlt]	53	1e-06
Query= sid 114859 lan dp1ORF038 Phage dp1 ORF 1350-1871 3 (173 letters)		
gi 1175542 sp P44123 YB90_HAEIN HYPOTHETICAL PROTEIN HI1190 >gi ...	100	6e-21
gi 2982977 (AE000681) hypothetical protein [Aquifex aeolicus]	67	7e-11
gi 3860744 emb CAA14645 (AJ235270) unknown [Rickettsia prowazekii]	65	3e-10
gi 2650193 (AE001074) conserved hypothetical protein [Archaeoglo...	58	4e-08
gi 3258383 dbj BAA31066.1 (AP000007) 157aa long hypothetical pr...	55	2e-07
gi 1001713 dbj BAA10550 (D64004) hypothetical protein [Synechoc...	50	8e-06
gi 4155434 (AE001516) putative (Helicobacter pylori J99]	50	1e-05
Query= sid 114860 lan dp1ORF039 Phage dp1 ORF 3306-3803 3 (165 letters)		
gi 1922884 emb CAA68244 (X99978) ORF7; hydrophobic protein [Lact...	64	5e-10
Query= sid 114862 lan dp1ORF041 Phage dp1 ORF 8208-8699 3 (163 letters)		
gi 2522313 (AF012906) dUTPase homolog [Bacillus subtilis] >gi 26...	108	2e-23
gi 2634150 emb CAB13650 (Z99113) similar to deoxyuridine 5'-tri...	108	3e-23
gi 3913546 sp O54134 DUT_STRCO DEOXYURIDINE 5'-TRIPHOSPHATE NUCL...	56	2e-07
gi 3913542 sp O48500 DUT_BPT5 DEOXYURIDINE 5'-TRIPHOSPHATE NUCLE...	52	3e-06
gi 3913548 sp O68992 DUT_CHLTE DEOXYURIDINE 5'-TRIPHOSPHATE NUCL...	50	1e-05
Query= sid 114867 lan dp1ORF046 Phage dp1 ORF 42774-43202 3 (142 letters)		
gi 1934764 emb CAB07984 (Z93946) hypothetical protein [bacterio...	287	2e-77
Query= sid 114901 lan dp1ORF080 Phage dp1 ORF 42490-42759 1 (89 letters)		
gi 1934763 emb CAB07983 (Z93946) hypothetical protein [bacterio...	147	1e-35
Query= sid 114912 lan dp1ORF091 Phage dp1 ORF 43189-43413 1 (74 letters)		
gi 1934765 emb CAB07985 (Z93946) holin [bacteriophage Dp-1]	63	2e-10

Table 32

Sequence of Dp1 published by Sheehan and al.. 4731 nucleotides.

1	tttaaatttt	ttgacaaagt	taattcaaat	tgtaccgctg	aagcaatttt	ccatgtatcc	actcaaagtt
71	gttcagtggtg	gtccaatcat	attaaaaatcg	aacttggttaa	tatctctact	ccttttagtg	aagcagagga
141	agaccttaaa	tatcgaattg	actcaaaagc	cgatcaaaag	ctaactaacc	aacagttgac	ggcactcacg
211	gaaaaggctc	aactacatga	cgcagaactg	aaagctaagg	ctacaatgga	gcagtttaagt	aacttagaaa
281	aggcttaatga	aggtagaatg	aaagctaagt	aagaagctat	caacaaatcg	gaacccgacc	taactctagc
351	ggcaagtcca	attgaagcta	ctatccaaga	acttgccggg	ctacgggaac	tgaagaagtt	cgctcgacagt
421	tgcagtagct	cttctaatac	aggcttaatt	atcggttaaga	acgacggtag	ctctaccatt	aaggatcaaa
491	gtgaccgaat	ttctatgttc	tccgcaggga	atgaagttat	gtaccttacc	caagggttca	ttcacatcga
561	taacgggatac	tttacccaat	ccattccaagt	cgcccgattt	agaacgggaac	aatactcggt	taatccagac
631	atgaacgtga	ttcggtagtg	aggataagga	gaataacatg	acaaaattta	tcaactcata	cggccctctt
701	cacttgaacc	tttacgtcga	acaagttagt	caggacgtaa	cgaacaactc	ctcgcgagtt	agttggcgag
771	ctactgtcga	cgcgatgga	gcttatcgaa	cggtggactta	tggaaatatt	agtaaccttt	ccgtatgggt
841	aaatgggtca	agtgttcata	gcagtcaccc	agactacgac	acgtccggcg	aagaggttaac	gctcgcaagt
911	ggagaagtga	ctgttctcca	caatagttag	gggacaaaga	caatgtccgt	ttgggcttcg	tttgacctta
981	ataacggcgt	tcacggaaat	atcactatct	ctactaatta	cactttagac	agtattccaa	ggctcacaca
1051	gatttctagt	tttgaggga	atcgaaatct	aggatcttta	catacgggta	tctttaaccg	aaaagtgaac
1121	tcttttaccg	atcaagtttg	gtaccgagtt	ttcggtagcg	actggataga	tttaggttaag	aaccatacta
1191	ctacgctatc	ctttacgccc	tcaactggat	tagcaaggtta	cttacctaaa	tcaagttccg	gaacaatcga
1261	catctgtatt	cgaaacctata	acggaaactac	gcaaatgggt	agtgcagctc	attcaaacgg	atggaggttc
1331	aacatccccg	attcagtagc	tcctactttt	tcgggcattt	ctttagtaga	cacgacttca	gcggttcgac
1401	agatttttaac	agggaacaac	ttccctccaa	tcactgtcga	cattcaagtc	aacttcaaca	atgcttccgg
1471	cgcttacgga	tccactatcc	aagcatttca	cgctgagctc	gtaggtaaaa	accaagctat	caacgaaaac
1541	ggcggcgaat	tgggtatgat	gaactttaat	ggctccgcta	ccgtaagagc	atgggttaca	gacacgcgag
1611	gaaaacaatc	gaacgtccaa	gacgtatcta	tcaatgttat	agaatactat	ggaccgtcta	tcaatttctc
1681	cggtccaacgt	actcgtaaaa	atccgtgcaat	tatccaagct	cttcgaaatg	ctaaggtcgc	acctataaag
1751	gtaggagggtc	aacagaaaaa	catcatgcaa	attacctctc	ccgtggcgcc	gttgaaactc	actaatttca
1821	cagaagatag	agggttcggcg	tcaggggagct	tcactactat	ttccctactg	actaactcgt	ccgcgaactt
1891	agctcggtaac	tacgggcccgg	acaagttctta	catagtttaag	gctaaaatcc	aagacaggtt	cacttcgact
1961	gaatttagtg	ctacggtagc	taccgaatca	gtagttctta	actatgacaa	ggacggctga	cttgagggtg
2031	gtaaggttgt	agaacaaggg	aaggcagggg	caattgatgc	agcaggtgat	atatatgctg	gaggtcgaca
2101	agttcaacag	tttcagctca	ctgataataa	tggagcattg	aacagggggc	aatataacga	tgttggaata
2171	agcgtgaaac	agagtttaca	tggcgaagta	acaaatacga	ggacaaccct	acgggaactc	gaggtgaatg
2241	gggactatct	caaaatttct	gggttagatg	ctggaaaatg	gttcaatcct	tcattacaat	gtcaggaaga
2311	atgttcatca	ggacagcgaa	cgatggaaac	agctggagac	ctaacaagtg	gaaagaggtt	ctattttaagc
2381	aagacttcga	acagaataat	tggcagaaac	ttgttcttca	aagtgggtgg	aacctacact	caacctatgg
2451	cgacgcattc	tattcgaaaa	ctcttgacgg	catagtatat	ttgagaggaa	atgtgcataa	aggacttatc
2521	gacaaaaggag	ctactattgc	agtacttctc	gaaggattta	gaccgaaagt	ttcaatgtat	ctatcttagc
2591	tcaataactc	atatggaaat	gccattctat	gtatatacac	tgacgggaag	cttgtggtga	aatcgaatgt
2661	agataattct	tgggttaaat	tagacaatgt	ctcatttcgt	atttaatttg	agctgaaatc	atgataatgt
2731	attttttaga	aaaggagggtg	gaactatgtt	gaaccttaca	aaatcgcgcc	aaattgtggc	agagttcact
2801	attgggcaag	gagctgaaaa	gaaactgttc	aaaacaacga	ttgtgaacat	tgatgcaaac	gcagtatcaa
2871	ccgtctctga	aactcttcat	gaccagactc	tgtatgctgc	gaaccgtcga	gaacttcgag	ctgacgagca
2941	aaaacttcgc	gaaactcggt	acgcaatcga	agatgaaatt	aatagctgga	gcggggggaa	aaagggggag
3011	cccgggtcta	acaggctgaa	taaggaggcg	tcaatctatg	ccaatgtggc	taaacgacac	cgcagctctg
3081	acgacgatta	ttacagcgtg	cagcggagtg	cttactgtcc	tactaaataa	gttatttcgaa	tggaaatcga
3151	ataaaggcaa	gagcgtttta	gaggatattc	ctacaactct	tagcactctt	aaacagcagg	tcgacgggat
3221	tgaccaaacg	acagtagcaa	tcaatcacca	aaatgacgtc	attcaagacg	gaactagaaa	aattcaacgt
3291	taccgtcttt	atcacgactt	aaaaaggga	gtgataacag	gctatacaac	tctcgaccat	tttagagagc
3361	tctctatttt	attcgaaagt	tataaagaac	tggcgggaaa	tgggtgaagt	gaagccttgt	atgaaaaata
3431	caagaaatta	ccaattaggg	aggaagattt	agatgaaact	atctaacgaa	caatatgacg	tagcaaaaga
3501	cggtggaacc	gtagtcgttc	cagcagcgat	tgacttaatt	acaggtcttg	gagcgttgta	tcaatttgac
3571	actactgcta	tcacagggaac	cattgcactt	cttgcaactt	ttgcagggtac	tgcttctagga	gtttctagcc
3641	gaaactacca	aaaggaaaca	gaagctcaaa	acaatgaggt	ggaataatgg	gagtcgatat	tgaaaaaggc
3711	gttcgctgga	tgaggcccg	aaagggtcga	gtatcttata	gcatggactt	tcgagacggt	cctgatagct
3781	atgactgctc	aagttctatg	tactatgctc	tccgctcagc	cggagcttca	agtgctggat	gggcagtcga
3851	tactgagtag	atgcacgcac	ggcttattga	aaacgggtat	gaactaatta	gtgaaaatgc	tccgtgggat
3921	gctaaacgag	gcgacatctt	catctgggga	cgcaaaagggtg	ctagcgcagg	cgctggagggt	catacagggg
3991	tggttcattga	cagtgataac	atcattcact	gcaactacgc	ctacgacgga	atttccgtca	acgaccacga
4061	tgagcgttgg	tactatgcag	gtcaacctta	ctactacgtc	tatcgcttga	ctaacgcaaa	tgctcaaccg
4131	gctgagaaga	aacttggtg	gcagaaagat	gctactgggt	tctggtacgc	tcgagcaaac	ggaaacttatc
4201	caaaagatga	gttcgagtat	atcgaaagaa	acaagttctg	gttctacttt	gacgaccaag	gctacatgct
4271	cgctgagaaa	tgggtgaaac	atactgatgg	aaattgggtat	tgggttcgac	gtgacgggata	ctgggctacg
4341	tcatggaaac	ggattggcga	gtcactggatc	tacttcaatc	gcgatgggtt	aatggtaacc	gggttgattta
4411	agtattacga	taattgggtat	tattgtgatg	ctaccaacgg	cgacatgaaa	tcgaatcggt	ttatccggtta
4481	tacacgacgg	tggtatctac	tattaccgga	cggacgtctg	gcagataaac	ctcaattcac	cgtagagcga
4551	gacgggctca	ttactgctaa	agtttataat	atagagagga	ggaagctctt	ttcttaatat	tggttctctt
4621	aatcccgcaa	gggttcgacc	ctgcgggggt	tatgtgtcgt	gaattactct	atttacttat	tcgaagattt
4691	caattataat	taataaatca	acgagattca	taattggagg	aatg		

Table 33

Streptococcus accession numbers

gi 5776553 gb AF026471.2 AF026471 [5776553]	gi 5231200 gb AF157824.1 AF157824 [5231200]
gi 5410470 gb AF139890.1 AF139890 [5410470]	gi 5231197 gb AF157823.1 AF157823 [5231197]
gi 5410468 gb AF139889.1 AF139889 [5410468]	gi 5231194 gb AF157822.1 AF157822 [5231194]
gi 5410466 gb AF139888.1 AF139888 [5410466]	gi 5231191 gb AF157821.1 AF157821 [5231191]
gi 5410464 gb AF139887.1 AF139887 [5410464]	gi 5231188 gb AF157820.1 AF157820 [5231188]
gi 5410462 gb AF139886.1 AF139886 [5410462]	gi 5231185 gb AF157819.1 AF157819 [5231185]
gi 5410460 gb AF139885.1 AF139885 [5410460]	gi 5231182 gb AF157818.1 AF157818 [5231182]
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CLAIMS

What is claimed is:

- 5 1. A method for identifying a bacteriophage coding region encoding a product active on an essential bacterial target, comprising identifying a nucleic acid sequence encoding a gene product which provides a bacteria-inhibiting function when said bacteriophage infects a host bacterium,
- 10 wherein said bacteriophage is uncharacterized and said host bacterium is a pathogenic bacterium.
- 15 2. The method of claim 1, further comprising expressing a recombinant bacteriophage ORF in cells of a bacterial strain, wherein inhibition of said cells following expression of said ORF is indicative that said product is active on an essential bacterial target.
- 20 3. The method of claim 2, wherein inhibition of said bacterium following expression of said ORF is determined by comparison with the growth or viability of said bacterium following expression of an inactivated mutant form of said ORF or in the absence of expression of said ORF, and wherein inhibition of said bacterium following expression of said ORF is indicative that said product is active on an essential bacterial target.
- 25 4. The method of claim 2, wherein expression of said ORF is inducible.
5. The method of claim 1, further comprising sequencing at least a portion of a bacteriophage genome.
- 30 6. The method of claim 1, wherein at least a portion of the nucleotide sequence of a bacteriophage genome is known, said method further comprising identifying at least one ORF in said portion by computer analysis of said sequence.
- 35 7. The method of claim 6, further comprising analyzing the sequence of said at least one ORF or of a polypeptide encoded by said ORF to identify homologous genes or gene products of known biochemical function, thereby indicating the biochemical function of said polypeptide.

8. The method of claim 7, wherein said homologous gene or gene product is a bacterial gene important for cell viability.

9. The method of claim 7, wherein said homologous gene or gene product is a gene or gene product known to have a bacteria-inhibiting function.

10. The method of claim 6, further comprising analyzing the sequence of said at least one ORF or of a polypeptide encoded by said ORF to identify structural motifs in said polypeptide, thereby indicating the cellular function of said polypeptide.

11. The method of claim 1, wherein a host bacterium for said bacteriophage is selected from the species group consisting of bacteria listed in Table 1.

12. The method of claim 1, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

13. The method of claim 2, wherein a plurality of bacteriophage ORFs are expressed in at least one bacterium.

14. The method of claim 13, wherein each of said plurality of bacteriophage ORFs is expressed in a different bacterium.

15. The method of claim 14, wherein said plurality of bacteriophage ORFs comprises at least 10% of the ORFs in the genome of said bacteriophage.

16. The method of claim 1, wherein said pathogenic bacterium is an animal pathogen.

17. The method of claim 16, wherein said pathogenic bacterium is a human pathogen.

18. The method of claim 1, wherein said pathogenic bacterium is a plant pathogen.

19. The method of claim 1, further comprising confirming the inhibitor function of said ORF.

20. The method of claim 19, wherein said confirming comprises expressing a loss-of-function mutant form of said ORF in said host bacterium.

5 21. The method of claim 1, wherein said identifying a nucleic acid sequence encoding a gene product active on an essential bacterial target comprises identifying a nucleic acid sequence encoding a homolog of a bacteriophage polypeptide known to be active on an essential bacterial target.

10 22. The method of claim 1, wherein said identifying a bacteriophage coding region comprises identifying a first coding region from a bacteriophage having a non-pathogenic host bacterial strain related to said pathogenic bacterium, said first coding region encoding a product active on an essential bacterial target; and
identifying a homolog of said first coding region, wherein said
15 homolog is a probable said bacteriophage coding region encoding a product active on an essential bacterial target.

23. The method of claim 2, wherein a plurality of bacteriophage ORFs from a plurality of different bacteriophage are expressed in at least one bacterium.
20

24. The method of claim 23, wherein each of said plurality of bacteriophage ORFs are expressed in different bacteria.

25 25. A method for identifying a target for antibacterial agents, comprising determining the bacterial target of an uncharacterized bacteriophage inhibitor protein.

26. The method of claim 25, wherein said determining comprises identifying at least one bacterial protein which binds to said bacteriophage inhibitor
30 protein or a fragment thereof.

27. The method of claim 26, wherein said binding is determined using affinity chromatography on a solid matrix.

35 28. The method of claim 25, wherein said determining comprises identifying at least one protein:protein interaction using a genetic screen.

29. The method of claim 28, wherein said genetic screen is a yeast two-hybrid screen.

30. The method of claim 25, wherein said determining comprises a co-immunoprecipitation assay or a protein-protein crosslinking assay.

31. The method of claim 25, wherein said determining comprises identifying a mutated bacterial coding sequence which protects a bacterium from said bacteriophage inhibitor.

10

32. The method of claim 25, wherein said determining comprises identifying a bacterial coding sequence which protects a bacterium against said bacteriophage inhibitor when expressed at high levels in said bacterium.

33. The method of claim 25, wherein said determining further comprises identifying a bacterial nucleic acid sequence encoding a polypeptide target of said bacteriophage inhibitor protein.

15

34. The method of claim 33, wherein said nucleic acid sequence is identified by determining at least a portion of the amino acid sequence of a bacterial protein target, and identifying a bacterial nucleic acid sequence which encodes said protein target.

20

35. The method of claim 25, wherein said bacterial target is naturally produced by a bacterial species selected from the group consisting of species of the genera listed in Table 1.

25

36. The method of claim 25, wherein said bacterial target is naturally produced by a bacterial strain selected from the group consisting of species listed in Table 1.

30

37. The method of claim 25, wherein said inhibitor protein is naturally produced by a bacteriophage selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

35

38. The method of claim 25, further comprising identifying a bacteriophage ORF which encodes a product having a bacteria-inhibiting function.

39. The method of claim 38, wherein said identifying a phage ORF comprises expressing at least one bacteriophage ORF in a bacterium, wherein inhibition of said bacterium following said expression is indicative that said ORF
5 encodes a bacteria-inhibiting function.
40. The method of claim 39, wherein a plurality of bacteriophage ORFs are expressed in at least one bacterium.
- 10 41. The method of claim 40, wherein each of said plurality of bacteriophage ORFs is expressed in a different bacterium.
42. The method of claim 41, wherein said plurality of bacteriophage ORFs comprises at least 10% of the ORFs in the genome of said bacteriophage.
15
43. The method of claim 25, wherein said determining the bacterial target of a bacteriophage inhibitor protein is performed for a plurality of different bacteriophage of the same host bacterium.
- 20 44. The method of claim 25, wherein said bacterial target originates from an animal pathogen.
45. The method of claim 44, wherein said bacterial target is a gene homologous to a gene from an animal pathogen.
25
46. The method of claim 44, wherein said pathogen is a human pathogen.
47. The method of claim 25, wherein said bacterial target originates from a plant pathogen.
30
48. The method of claim 25, wherein said bacterial target is a gene homologous to a gene from a plant pathogen.
49. The method of claim 25, further comprising determining the cellular or
35 biochemical function or both of said inhibitor protein.

50. The method of claim 25, wherein said identifying the bacterial target comprises identifying a phage-specific site of action.

5 51. An isolated, purified, or enriched nucleic acid sequence at least 15 nucleotides in length, wherein said sequence corresponds to at least a portion of a bacteriophage sequence, and wherein said bacteriophage is selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD, *Enterococcus* bacteriophage 182, and *Streptococcus pneumoniae* bacteriophage Dp-1.

10

52. The nucleic acid sequence of claim 51, wherein said sequence comprises at least 50 nucleotides.

15 53. The nucleic acid sequence of claim 51, wherein said nucleic acid sequence corresponds to at least a portion of a nucleic acid sequence which encodes a product which provides a bacteria-inhibiting function.

20 54. The nucleic acid sequence of claim 53, wherein said nucleic acid sequence encodes a polypeptide which provides a bacteria-inhibiting function.

55. The nucleic acid sequence of claim 54, wherein said nucleic acid sequence is transcriptionally linked with regulatory sequences enabling induction of expression of said sequence.

25

56. An isolated, purified, or enriched polypeptide comprising at least a portion of a protein providing a bacteria-inhibiting function, wherein said polypeptide is normally encoded by a bacteriophage selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD, *Enterococcus* bacteriophage 182, and *Streptococcus pneumoniae* bacteriophage Dp-1.

30

57. The polypeptide of claim 56, wherein said polypeptide provides said bacteria-inhibiting function.

35 58. The polypeptide of claim 56, wherein said polypeptide comprises a portion at least 10 amino acid residues in length of a said polypeptide normally encoded by said bacteriophage.

59. A recombinant vector comprising a bacteriophage ORF corresponding to an ORF from a bacteriophage having a pathogenic bacterial host, wherein said
5 bacterial host is selected from the group consisting of uncharacterized bacteria of Table 1.

60. The vector of claim 59, wherein said vector is an expression vector.

10 61. The vector of claim 59, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage of Table 1.

62. The vector of claim 61, wherein said bacteriophage is selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD,
15 *Enterococcus* bacteriophage 182, and *Streptococcus pneumoniae* bacteriophage Dp-1.

63. The vector of claim 60, wherein expression of said ORF is inducible.

20 64. A recombinant cell comprising a vector, wherein said vector comprises an ORF from a bacteriophage having a pathogenic bacterial host, wherein said bacterial host is selected from the group consisting of bacterial species of Table 1.

65. The recombinant cell of claim 64, wherein said bacteriophage is
25 selected from the group consisting of uncharacterized phage of Table 1.

66. The cell of claim 65, wherein said bacteriophage is selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD,
30 *Enterococcus* bacteriophage 182, and *Streptococcus pneumoniae* bacteriophage Dp-1.

67. The cell of claim 64, wherein said vector is an expression vector and expression of said ORF is inducible.

35 68. A method for identifying an antibacterial agent, comprising identifying an active portion of a product of a bacteria-inhibiting ORF of a bacteriophage.

69. The method of claim 68, further comprising constructing a synthetic peptidomimetic molecule, wherein the structure of said molecule corresponds to the structure of said active portion.

5

70. A method for identifying a compound active on a target of a bacteriophage inhibitor protein, comprising the step of contacting a bacterial target protein with a test compound; and determining whether said compound binds to or reduces the level of activity of said target protein, wherein binding of said compound with said target protein or a reduction of the level of activity of said protein is indicative that said compound is active on said target and wherein said target is uncharacterized.

15 71. The method of claim 70, wherein said contacting is carried out *in vitro*.

72. The method of claim 70, wherein said contacting is carried out *in vivo* in a cell.

20 73. The method of claim 70, wherein said compound is a small molecule.

74. The method of claim 70, wherein said compound is a peptidomimetic compound.

25 75. The method of claim 70, wherein said compound is a fragment of a bacteriophage inhibitor protein.

76. The method of claim 70, further comprising determining the site of action of said compound on said target protein.

30

77. The method of claim 70, wherein said contacting is performed for a plurality of said target proteins.

35 78. A method of screening for potential antibacterial agents, comprising the step of determining whether any of a plurality of compounds is active on a target of a bacteriophage inhibitor protein,

wherein said target is naturally produced by a pathogenic bacterium.

79. The method of claim 78, wherein said plurality of compounds are small molecules.

5

80. The method of claim 78, wherein said determining is performed for a plurality of said targets.

10

81. A method for inhibiting a bacterium, comprising the step of; contacting said bacterium with a compound active on a target of a bacteriophage inhibitor protein, wherein said target or the target site is uncharacterized.

15

82. The method of claim 81, wherein said compound is said protein or an active fragment thereof.

83. The method of claim 81, wherein said compound is a structural mimetic of said protein.

20

84. The method of claim 81, wherein said compound is a small molecule.

85. The method of claim 81, wherein said contacting is performed *in vitro*.

25

86. The method of claim 81, wherein said contacting is performed *in vivo* in an animal.

87. The method of claim 86, wherein said animal is a human.

30

88. The method of claim 81, wherein said contacting is carried out *in vivo* in a plant.

89. The method of claim 81, wherein said bacterium is selected from the group of bacteria listed in Table 1.

35

90. A method for treating a bacterial infection in an animal suffering from an infection, comprising administering to said animal a therapeutically effective amount of compound active on a target of a bacteriophage inhibitor protein in a bacterium involved in said infection,
- 5 wherein said target is an uncharacterized target or the compound is active at an uncharacterized target site.
91. The method of claim 90, wherein said compound is a small molecule.
- 10 92. The method of claim 90, wherein said compound is a peptidomimetic compound.
93. The method of claim 90, wherein said compound is a fragment of a bacteriophage inhibitor protein.
- 15 94. The method of claim 90, wherein said animal is a mammal.
95. The method of claim 94, wherein said mammal is a human.
- 20 96. The method of claim 90, wherein said bacterium is selected from the group listed in Table 1.
97. The method of claim 90, wherein said bacteriophage inhibitor protein is from a bacteriophage selected from the group of bacteriophage listed in Table 1.
- 25 98. A method for prophylactically treating an animal at risk of an infection, comprising administering to said animal a prophylactically effective amount of a compound active on a target of a bacteriophage inhibitor protein,
- 30 wherein said target is an uncharacterized target or the site of action of said compound is an uncharacterized target site.
99. The method of claim 98, wherein said compound is a small molecule.
- 35 100. The method of claim 98, wherein said compound is a peptidomimetic compound.

101. The method of claim 98, wherein said compound is a fragment of a bacteriophage inhibitor protein.
102. The method of claim 98, wherein said animal is a mammal.
- 5 103. The method of claim 102, wherein said mammal is a human.
104. An antibacterial agent active on a target of a bacteriophage inhibitor protein, wherein said target is an uncharacterized target or said agent is active at a phage-specific site on said target.
- 10 105. The agent of claim 104, wherein said agent is a pepetidomimetic of a bacteriophage inhibitor polypeptide.
- 15 106. The agent of claim 104, wherein said agent is a small molecule.
107. The agent of claim 104, wherein said agent is a fragment of a bacteriophage inhibitor polypeptide.
- 20 108. The agent of claim 104, wherein said agent is active at a phage-specific site on said target.
- 25 109. A method of making an antibacterial agent, comprising the steps of:
- a) identifying a target of a bacteriophage inhibitor polypeptide;
 - b) screening a plurality of test compounds to identify a compound active on said target; and
 - c) synthesizing said compound in an amount sufficient to provide a
- 30 therapeutic effect when administered to an organism infected by a bacterium naturally producing said target.
110. The method of claim 109, wherein said compound is a small molecule.
- 35 111. The method of claim 109, wherein said compound is a peptidomimetic compound.

112. The method of claim 109, wherein said compound is a fragment or derivative of a bacteriophage inhibitor protein.

5 113. A computer readable device having recorded therein a nucleotide sequence of a portion of at least one bacteriophage genome of *Staphylococcus aureus* bacteriophage 77, bacteriophage 3A, or bacteriophage 96, a nucleotide sequence at least 95% identical to a said nucleotide sequence, a ribonucleic acid equivalent, a degenerate equivalent, a homologous sequence, or at least one amino acid sequence
10 encoded by said nucleotide sequence; and
 a nucleotide sequence or amino acid sequence analysis program,
 wherein said program can perform at least one sequence analysis on said nucleotide or amino acid sequence.

15 114. The device of claim 113, wherein said at least a portion of at least one bacteriophage genome comprises at least one ORF.

 115. The device of claim 113, wherein said device comprises a medium selected from the group consisting of floppy disk, computer hard drive, optical disk,
20 computer random access memory, and magnetic tape wherein said nucleotide or amino acid sequence or said program or both are recorded on said medium.

 116. The device of claim 113, wherein said portion of at least one bacteriophage genomic nucleotide sequence comprises at least 50% of at least one
25 bacteriophage genomic sequence.

 117. The device of claim 113, wherein said at least one bacteriophage nucleotide genomic sequence comprises portions of a plurality of bacteriophage nucleotide genomic sequences.
30

 118. A computer-based system for identifying biologically important portions of a bacteriophage genome, comprising:
 a) a data storage medium having recorded thereon a nucleotide sequence
35 corresponding to a portion of at least one bacteriophage genome, wherein said bacteriophage genome is uncharacterized;

- b) a set of instructions allowing searching of said sequence to analyze said sequence; and
- c) an output device.

5 119. The system of claim 118, wherein said output device comprises comprises a device selected from the group consisting of a printer, a video display, and a recording medium.

10 120. The system of claim 118, wherein said bacteriophage genome is of a bacteriophage selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

15 121. The system of claim 118, wherein said uncharacterized bacteriophage is selected from the group consisting of bacteriophage 77, 3A, and 96.

 122. A method for identifying or characterizing a bacteriophage ORF, comprising the steps of:

- 20 a) providing a computer-based system for analyzing nucleic acid or amino acid sequence data, wherein said system comprises a data storage medium having recorded thereon at least one nucleotide or amino acid sequence corresponding to a portion of at least one uncharacterized bacteriophage genome, a set of instructions allowing searching of said sequence to analyze said sequence; and an output device;
- b) analyzing at least a portion of at least one said sequence; and
- 25 c) outputting results of said analyzing to said output device.

 123. The method of claim 122, wherein said analysis identifies sequence similarity or homology with sequences selected from the group consisting of bacterial ORFs encoding products with related biological function; ORFs encoding known
30 inhibitors or bacteria, essential bacterial ORFs.

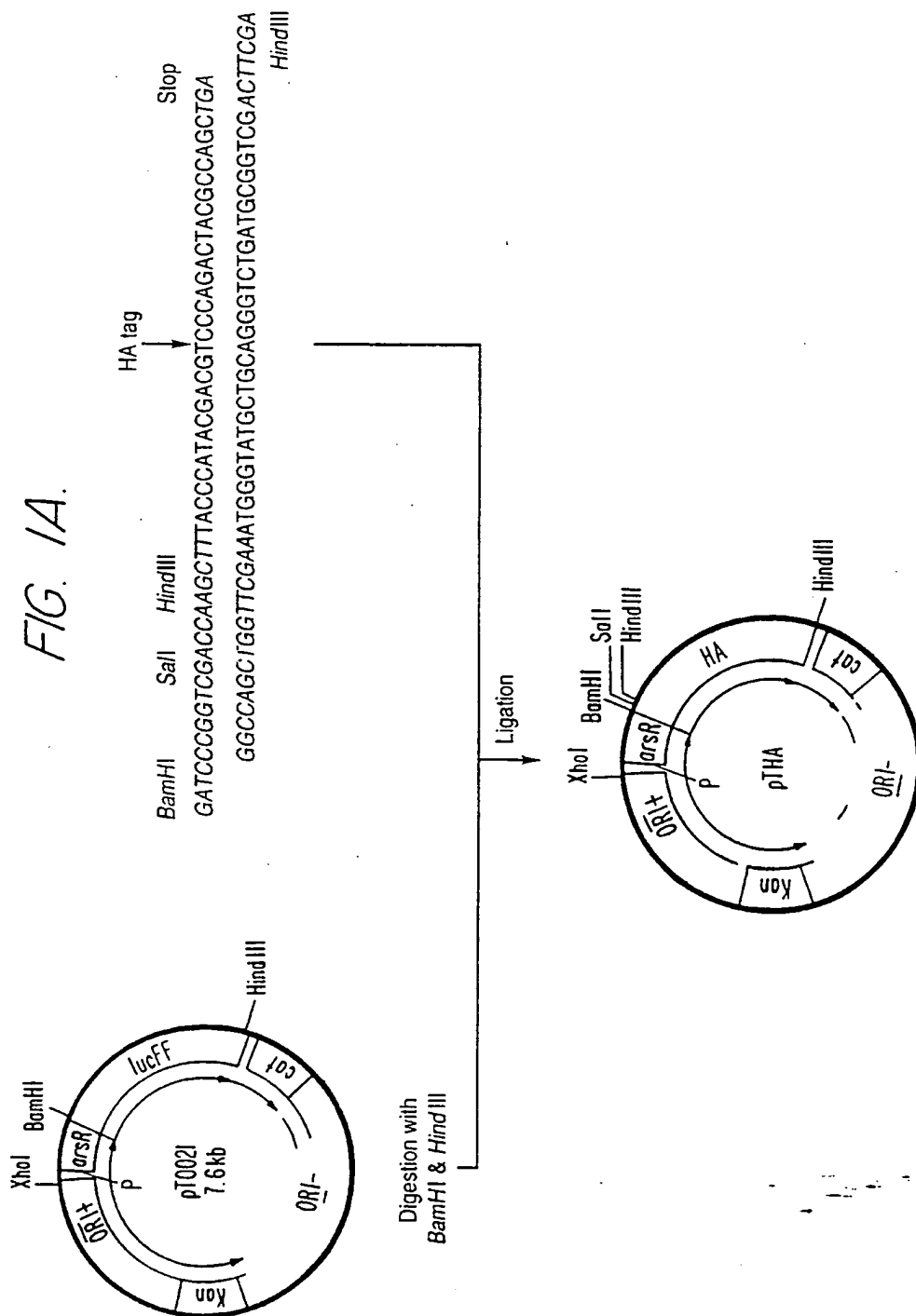
 124. The method of claim 122, wherein said analysis comprises identifying a probable biological function based on identification of structural elements or sequence homology or similarity.

35 125. The method of claim 122, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

126. The method of claim 125, wherein said uncharacterized bacteriophage is selected from bacteriophage 77, 3A, and 96.

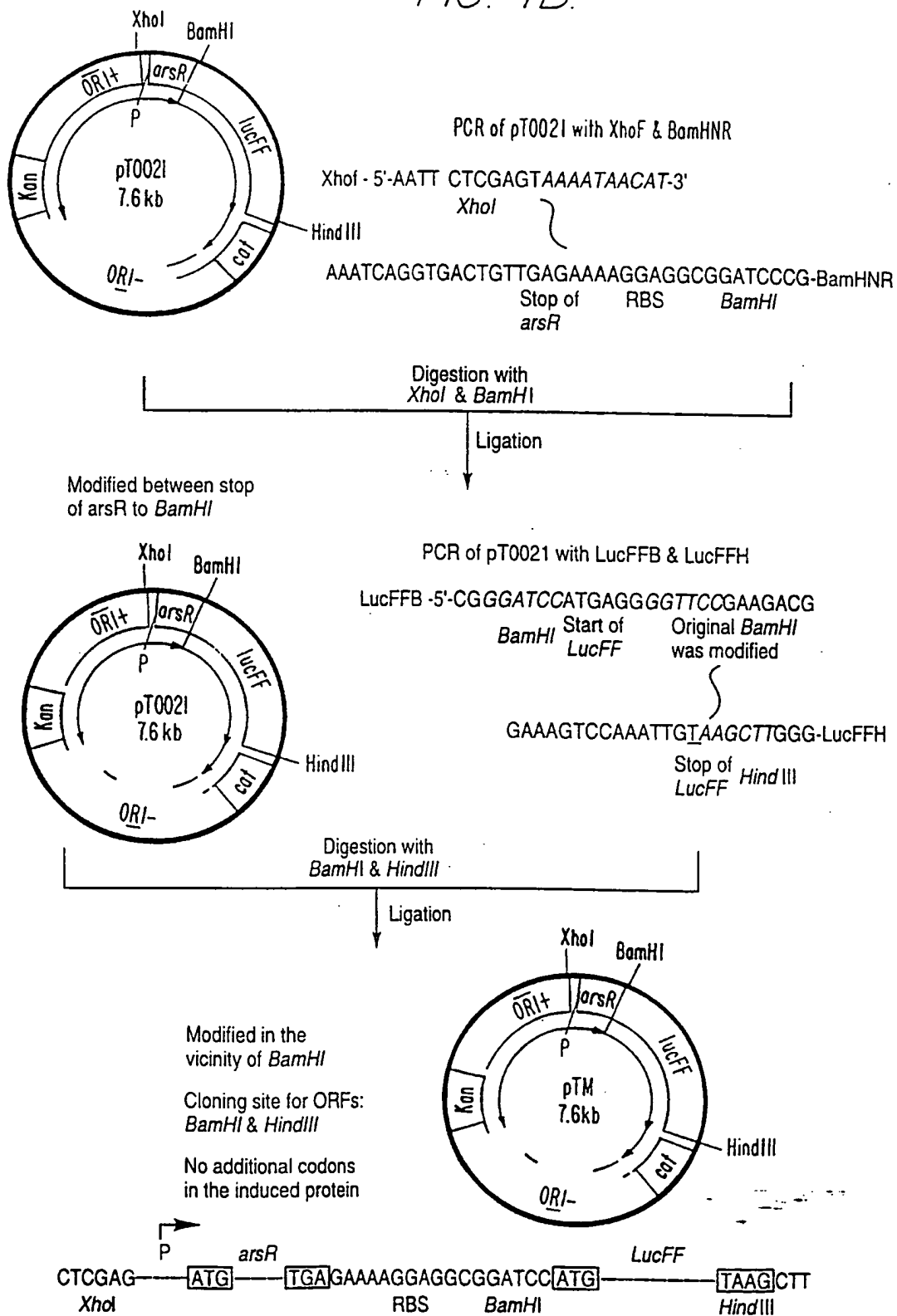
01/11

FIG. 1A.



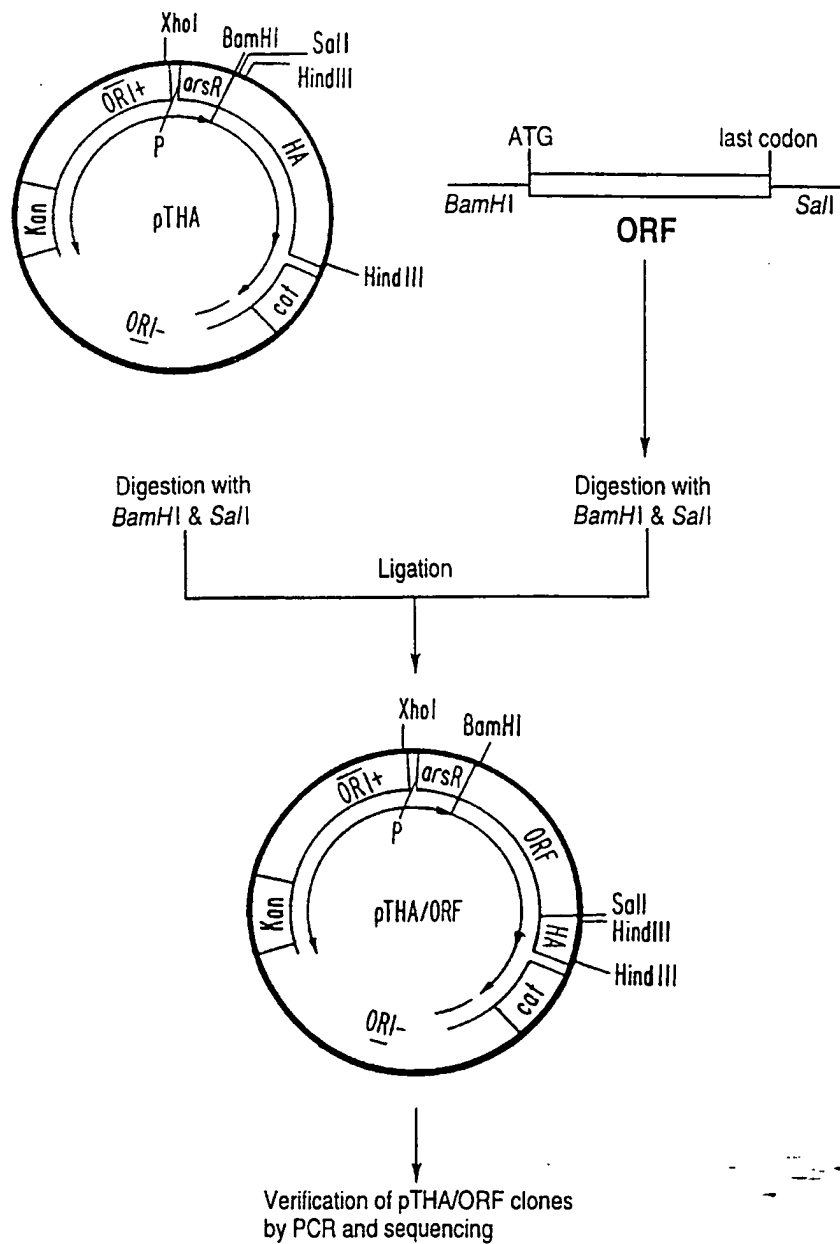
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FIG. 1B.



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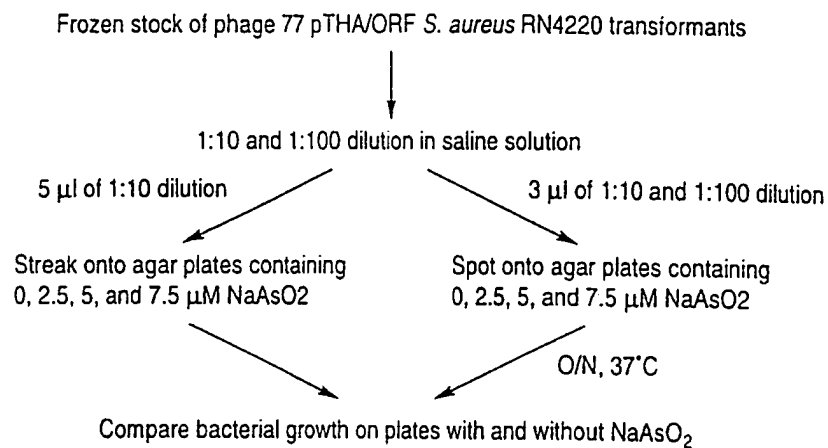
FIG. 2.



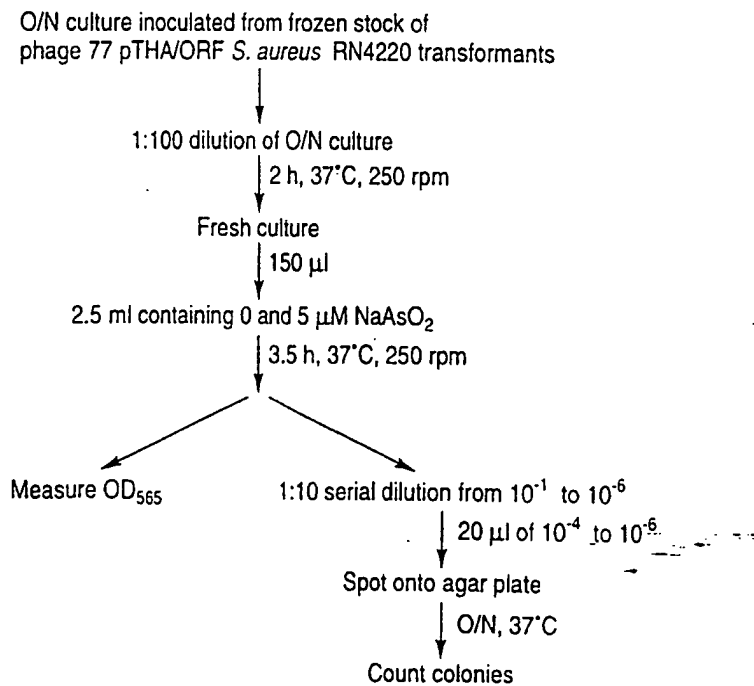
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FIG. 3.

(A) Functional assay on semi-solid support media



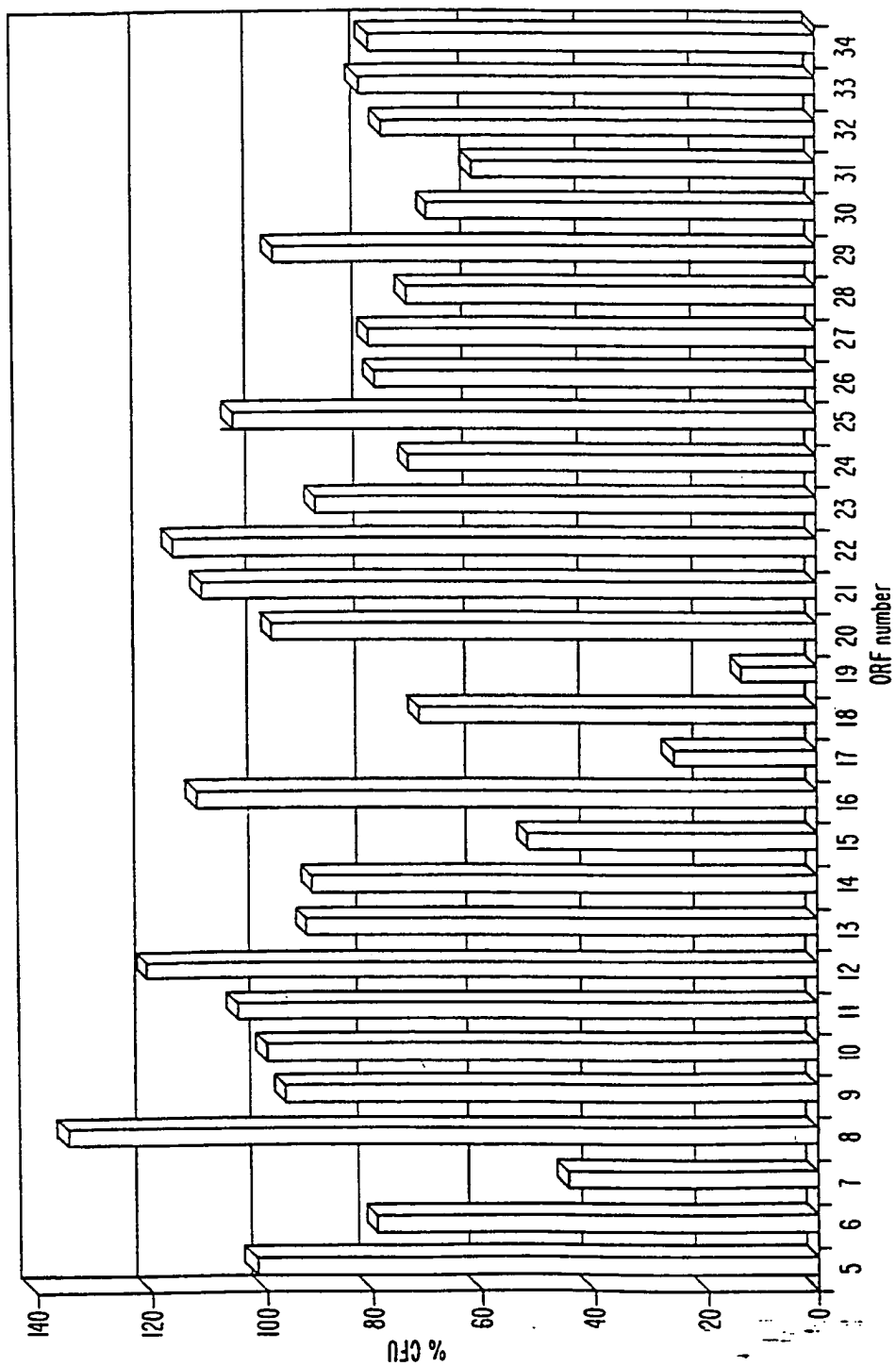
(B) Functional assay in liquid medium



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A. Inhibition of bacterial growth with individual ORFs of a *S. aureus* Bacteriophage.

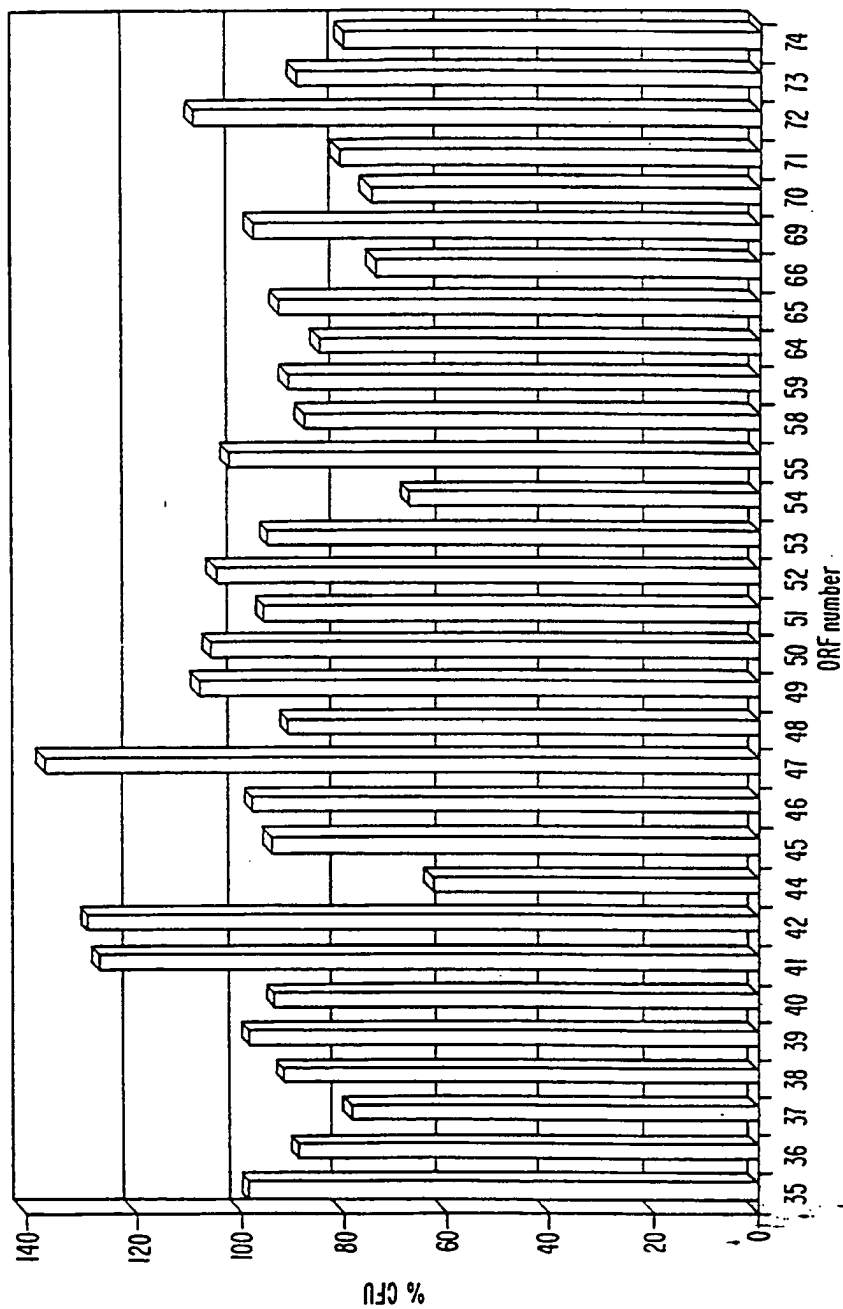
FIG. 4A.



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B. Inhibition of bacterial growth with individual ORFs of a *S. aureus* Bacteriophage.

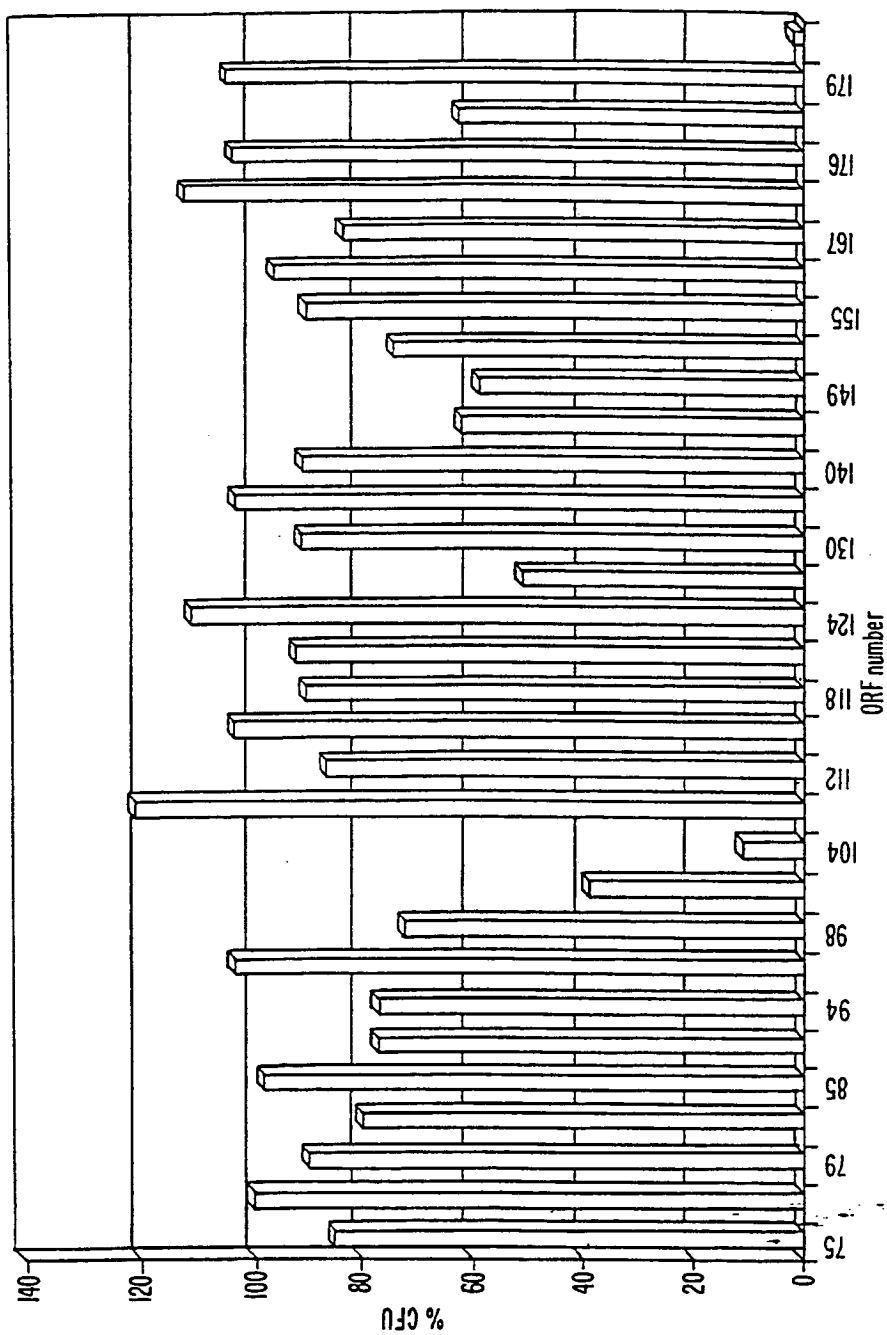
FIG. 4B.



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C. Inhibition of bacterial growth with individual ORFs of a *S. aureus* Bacteriophage.

FIG. 4C



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FIG. 5.

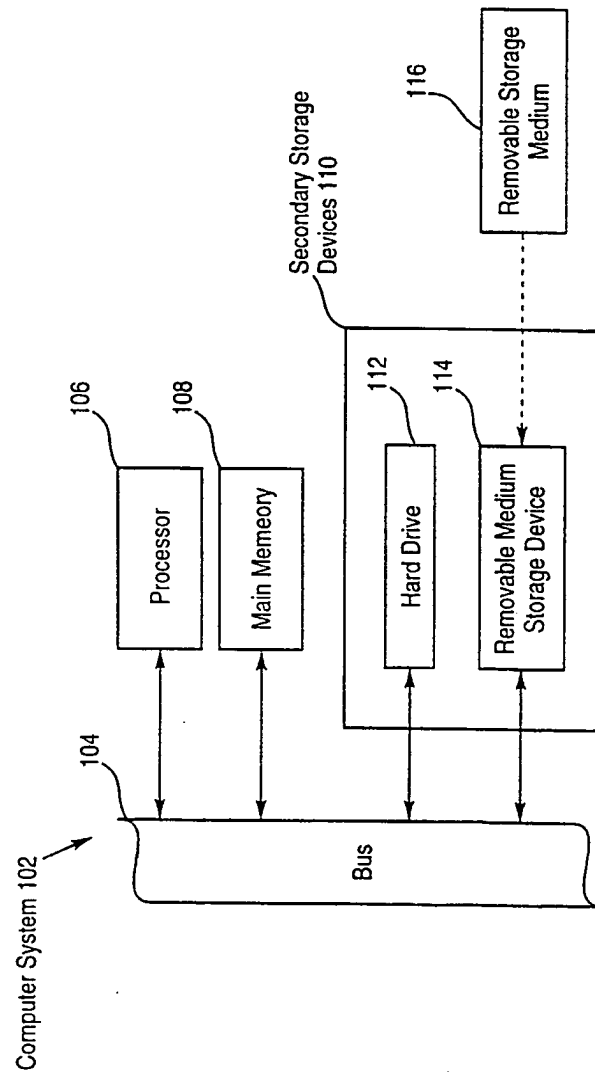
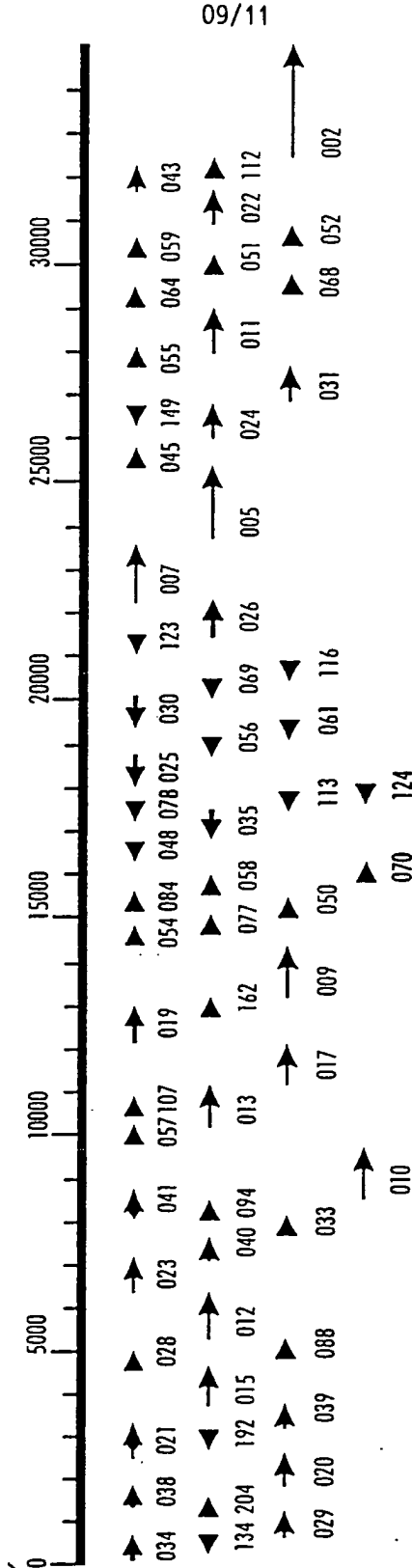


Fig 6A Fig 6B

Fig. 6

Fig. 6A

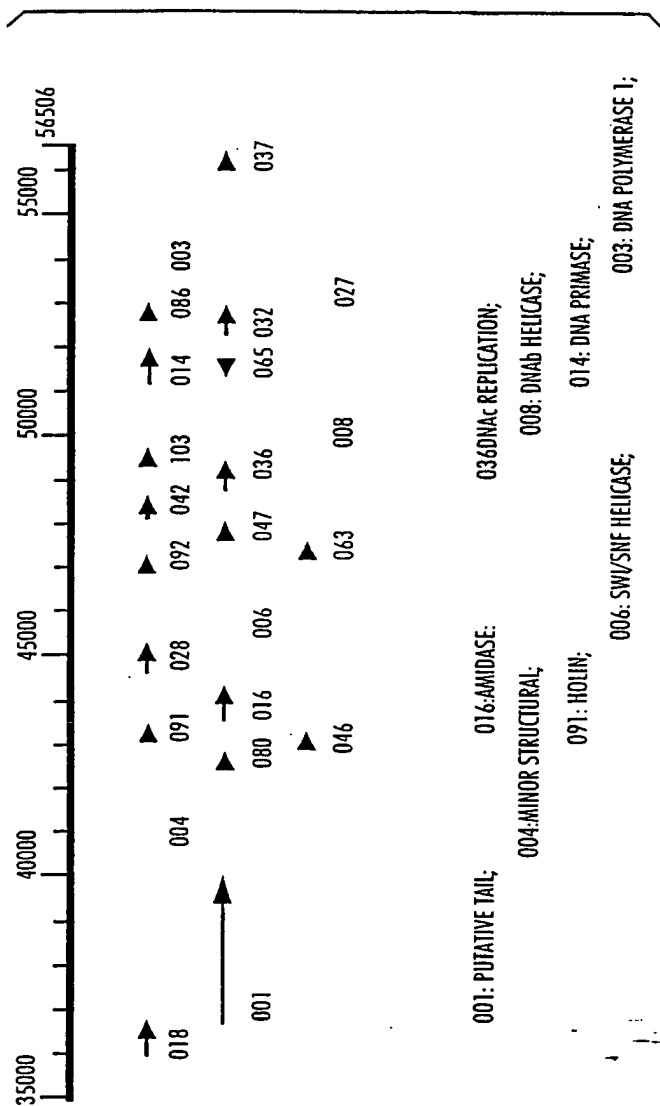
PHAGE: BACTERIOPHAGE Dp1
MINIMAL ORF SIZE: 33 A. A.
ORFS "WITH" RBS.
NUMBER OF ORFS: 85



029: exsB; 012: DNA pol. III beta;
038: exsC; 6-PYRUVYLTERAHDROPTERIN
020: exsD; COENZYME PQQ; 013: DNA POL. III GAMMA AND TAU;
021: GTP CYCLOHYDROLASE;
039: CITRULLINE BIOSYNTHESIS;
041: dUTPASE;
010: RecA;
007: TERMINASE;
011: MAJOR HEAD;
002: TAIL;

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Fig. 6B



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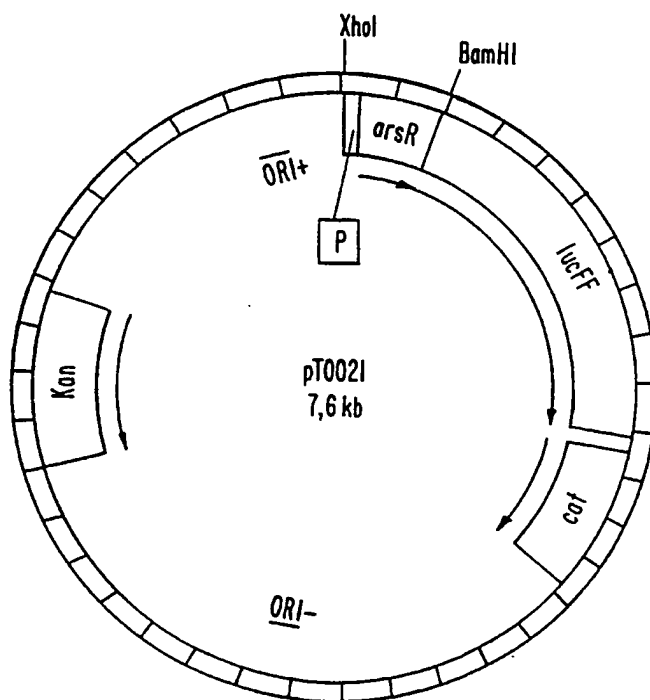
FIG. 7.

Abbreviations:

kan: gene encoding kanamycin resistance
 cat: gene encoding chloramphenicol resistance
 ori + and -: origin of replication in gram-positive and gram-negative bacteria, respectively
 arsR: gene encoding regulatory protein of the ars promoter
 P: ars promoter
 lucFF: gene encoding luciferase protein. This portion will be removed and replaced by individual *S. aureus* phage genes.

Reference:

Tauriainen et al., Appl. Environ. Microbio. 1997. 63: 4456-4461



(19) World Intellectual Property Organization
International Bureau



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60/157,218	30 September 1999 (30.09.1999)	US
60/168,777	1 December 1999 (01.12.1999)	US
09/454,252	2 December 1999 (02.12.1999)	US

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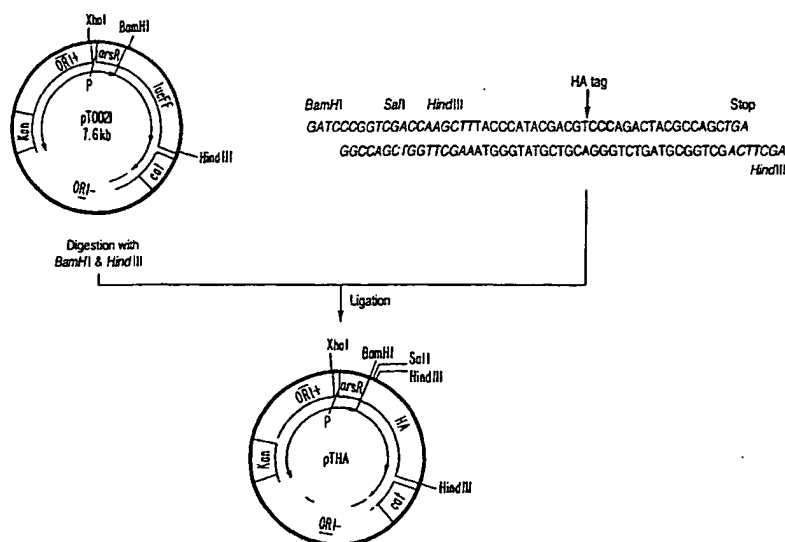
(72) Inventors; and

(75) Inventors/Applicants (*for US only*): PELLETIER,

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[Continued on next page]

(54) Title: DEVELOPMENT OF ANTI-MICROBIAL AGENTS BASED ON BACTERIOPHAGE GENOMICS



(57) Abstract: A method for identifying suitable targets for antibacterial agents based on identifying targets of bacteriophage-encoded proteins is described. Also described are compositions useful in the identification methods and in inhibiting bacterial growth, and methods for preparing and using such compositions.



Published:

— *With international search report.*

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

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INTERNATIONAL SEARCH REPORT

Intern. Application No

PCT/IB 99/02040

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C12Q1/70 C12Q1/68 C12N15/10 C12N15/34 C12N1/21
C07K14/01 C12Q1/18

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C12Q C12N C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 072 925 A (RUTGERS RES & EDUCATION FOUND) 2 March 1983 (1983-03-02)	1,2, 11-18
Y	the whole document	3-5,19, 20,22-24
X	--- SHEEHAN, M.M. ET AL.: "The lytic enzyme of the pneumococcal phage Dp-1: a chimeric lysin of intergeneric origin." MOLECULAR MICROBIOLOGY, vol. 25, no. 4, 1997, pages 717-25, XP000922620	1,2,11, 12,16,17
Y	the whole document	3-5,19, 20,22-24
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☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

3 July 2000

Date of mailing of the international search report

05.10.00

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INTERNATIONAL SEARCH REPORT

Inter. Appl. No.

PCT/IB 99/02040

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 95 27043 A (CARLTON RICHARD M ;MERRIL CARL R (US); US GOVERNMENT (US); ADHYA S) 12 October 1995 (1995-10-12) the whole document ---	1-3,5, 11,13, 16,17, 19,20
X	KANEKO J ET AL: "Complete nucleotide sequence and molecular characterization of the temperate staphylococcal bacteriophage phiPVL carrying Panton-Valentine leukocidin genes" GENE,NL,ELSEVIER BIOMEDICAL PRESS. AMSTERDAM, vol. 215, no. 1, pages 57-67, XP004149229 ISSN: 0378-1119 cited in the application the whole document ---	1,2,5, 11-13, 15-17, 22-24
X	WO 89 00199 A (UNIV LOUISIANA STATE) 12 January 1989 (1989-01-12) the whole document ---	1-3,5, 11-13, 15-17, 22-24
A	EP 0 748 871 A (NESTLE SA) 18 December 1996 (1996-12-18) the whole document -----	

INTERNATIONAL SEARCH REPORT

International application No.
PCT/IB 99/02040

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

invention 1: claims 2-4, 13-15, 23, 24 compl. and 1, 5, 11, 12, 16-20, 22, 68, 69 partially

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Invention 1: claims 2-4,13-15,23,24 completely,
and 1,5,11,12,16-20,22,68,69 partially

Method for identifying bacteriophage-encoded inhibitors of
pathogenic bacterial targets by expression of the viral ORF
in a host cell.

Invention 2: claims 6-10,21,
118-126 completely and 1,5,11,12,16-20,22,68,
69 partially

Method for identifying bacteriophage-encoded inhibitors of
pathogenic bacterial targets by computer-based methods, and
computer system for use therein.

Invention 3: claims 25-50

Method for identifying the pathogenic bacterial target of a
bacteriophage encoded product.

Invention 4: claims 51-67,81-103,113-117,
all partially, and as far as applicable

Isolated polynucleotides of at least 15 nucleotides in
length corresponding to at least a portion of sequence ID.1,
peptides comprising a portion of at least 10 amino acids
normally encoded by seq.ID.1, hosts, recombinant production
of the protein, computer-readable devices containing
sequence data of seq.ID.1, and a method for inhibiting a
pathogenic bacterium using the protein encoded by seq.ID.1,
all as far as applicable.

Inventions 5-2639 : claims 51-67,81-103,113-117,
all partially, and as far as applicable

Idem as invention 4, but limited to the respective seq.ID's
2-2636, whereby invention 5 relates to seq.ID.2, invention 6
relates to seq.ID.3,, and invention 2639 relates to
seq.ID.2636.

For the sake of conciseness, the general subject matter is
explicitly defined, the specific subject matters of each
invention are defined by analogy thereto.

Invention 2640: claims 70-80,109-112

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Method for identifying a compound active on the pathogenic bacterial target of a bacteriophage-encoded inhibitor and method for producing said compound.

Invention 2641: claims 103-108

Antibacterial agents active on the target of phage-encoded inhibitors in pathogenic bacteria.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/IB 99/02040

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
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